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(51) International Patent Classification 6:

WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(11) International Publication Number:

WO 98/39448

(43) International Publication Date: 11 September 1998 (11.09.98)

(30) Priority Data: 60/040,162 7 March 1997 (07.03.97) 60/040,333 7 March 1997 (07.03.97) 60/040,333 7 March 1997 (07.03.97) 60/040,161 7 March 1997 (07.03.97) 60/040,226 7 March 1997 (07.03.97) 60/040,334 7 March 1997 (07.03.97) 60/040,163 7 March 1997 (07.03.97) 60/040,163 7 March 1997 (07.03.97) 60/043,360 11 April 1997 (11.04.97) 60/043,568 11 April 1997 (11.04.97)	(22) International Filing Date: 6 March 1998 (06.03.98)	(21) International Application Number: PCT/US98/04493	C12N 15/12, 5/10, 1/21, C07K 14/47, 16/18, C12N 1/21, C07K 14/47, 16/18, C12Q 1/68, G01N 33/50, 33/53, 33/68, A61K 38/17
	06.03.9	98/0449	124
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> Without international search report and to be republished upon receipt of that report.

(54) Title: 186 HUMAN SECRETED PROTEINS

(57) Abstract

The present invention relates to 186 novel human secreted proteins and Isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.

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186 Human Secreted Proteins

Field of the Invention

encoded by these polynucleotides, uses of such polynucleotides and polypeptides, and This invention relates to newly identified polynucleotides and the polypeptides their production

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Background of the Invention

organelle, contains different proteins essential for the function of the organelle. The cell uses "sorting signals," which are amino acid motifs located within the protein, to target membrane, human cells and other eucaryotes are subdivided by membranes into many Unlike bacterium, which exist as a single compartment surrounded by a functionally distinct compartments. Each membrane-bounded compartment, or proteins to particular cellular organelles.

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One type of sorting signal, called a signal sequence, a signal peptide, or a leader another organelle called the Golgi apparatus. Here, the Golgi distributes the proteins to proteins. Once localized to the ER, both groups of proteins can be further directed to sequence, directs a class of proteins to an organelle called the endoplasmic reticulum vesicles, including secretory vesicles, the cell membrane, lysosomes, and the other (ER). The ER separates the membrane-bounded proteins from all other types of

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secretory granules) until exocytosis is triggered. Similarly, proteins residing on the cell proteins can fuse with the cell membrane and release their contents into the extracellular space - a process called exocytosis. Exocytosis can occur constitutively or after receipt of a triggering signal. In the latter case, the proteins are stored in secretory vesicles (or membrane can also be secreted into the extracellular space by proteolytic cleavage of a extracellular space as a secreted protein. For example, vesicles containing secreted Proteins targeted to the ER by a signal sequence can be released into the 'linker" holding the protein to the membrane. . 25

encoding human secreted proteins have been identified. These secreted proteins include and characterizing novel human secreted proteins and the genes that encode them. This knowledge will allow one to detect, to treat, and to prevent medical disorders by using Despite the great progress made in recent years, only a small number of genes pervasive role of secreted proteins in human physiology, a need exists for identifying the commercially valuable human insulin, interferon, Factor VIII, human growth hormone, tissue plasminogen activator, and erythropoeitin. Thus, in light of the secreted proteins or the genes that encode them.

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Summary of the Invention

polypeptides. Moreover, the present invention relates to vectors, host cells, antibodies, provided are diagnostic methods for detecting disorders related to the polypeptides, and and recombinant methods for producing the polypeptides and polynucleotides. Also therapeutic methods for treating such disorders. The invention further relates to The present invention relates to novel polynucleotides and the encoded screening methods for identifying binding partners of the polypeptides. S

Detailed Description

Definitions

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The following definitions are provided to facilitate understanding of certain terms used throughout this specification In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is contained within a cell, and still be "isolated" because that vector, composition of polynucleotide could be part of a vector or a composition of matter, or could be matter, or particular cell is not the original environment of the polynucleotide altered "by the hand of man" from its natural state. For example, an isolated 2

extracellular space, the secreted protein can undergo extracellular processing to produce In the present invention, a "secreted" protein refers to those proteins capable of signal sequence, as well as those proteins released into the extracellular space without being directed to the ER, secretory vesicles, or the extracellular space as a result of a necessarily containing a signal sequence. If the secreted protein is released into the a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage. 8 23

sequence contained in SEQ ID NO:X or the cDNA contained within the clone deposited coding region, with or without the signal sequence, the secreted protein coding region, with the ATCC. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the As used herein, a "polynucleotide" refers to a molecule having a nucleic acid Moreover, as used herein, a "polypeptide" refers to a molecule having the translated as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. amino acid sequence generated from the polynucleotide as broadly defined.

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In the present invention, the full length sequence identified as SEQ ID NO:X was often generated by overlapping sequences contained in multiple clones (contig

analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X was deposited with the American Type Culture Collection ("ATCC"). As shown in Table 1, each clone is identified by a cDNA Clone ID (Identifier) and the ATCC Deposit Number. The ATCC is located at 12301 Park Lawn Drive, Rockville Maryland 20852, USA. The ATCC deposit was made pursuant to the terms of the

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purposes of patent procedure.

Budapest Treaty on the international recognition of the deposit of microorganisms for

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, the complement thereof, or the cDNA contained within the clone deposited with the ATCC. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 μg/ml denatured, sheared salmon sperm DNA, followed by washing the filters in 0.1x SSC at about 65°C.

Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily accomplished through the manipulation of formamide concentration (lower percentages of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a

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pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA; followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH,PO4; 0.02M EDTA

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Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

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Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a

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complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

5 5 S or for other reasons. "Modified" bases include, for example, tritylated bases and stranded regions. In addition, the polynucleotide can be composed of triple-stranded double-stranded regions, hybrid molecules comprising DNA and RNA that may be or modified RNA or DNA. For example, polynucleotides can be composed of singlepolyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically unusual bases such as inosine. A variety of modifications can be made to DNA and contain one or more modified bases or DNA or RNA backbones modified for stability regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also single-stranded or, more typically, double-stranded or a mixture of single- and doubleregions, single- and double-stranded RNA, and RNA that is mixture of single- and and double-stranded DNA, DNA that is a mixture of single- and double-stranded modified forms. The polynucleotide of the present invention can be composed of any

ટ્ટ 20 35 3 as well as in a voluminous research literature. Modifications can occur anywhere in a to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and branched, for example, as a result of ubiquitination, and they may be cyclic, with or polypeptide, including the peptide backbone, the amino acid side-chains and the amino Such modifications are well described in basic texts and in more detailed monographs processing, or by chemical modification techniques which are well known in the art polypeptides may be modified by either natural processes, such as posttranslational may contain amino acids other than the 20 gene-encoded amino acids. The nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, given polypeptide may contain many types of modifications. Polypeptides may be or carboxyl termini. It will be appreciated that the same type of modification may be Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent present in the same or varying degrees at several sites in a given polypeptide. Also, a formation, demethylation, formation of covalent cross-links, formation of cysteine, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a from posttranslation natural processes or may be made by synthetic methods. without branching. Cyclic, branched, and branched cyclic polypeptides may result The polypeptide of the present invention can be composed of amino acids joined

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formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990);

"SEQ ID NO:X" refers to a polynucleotide sequence while "SEQ ID NO:Y" refers to a polypeptide sequence, both sequences identified by an integer specified in Table 1.

Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

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"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency dose exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.)

25 Polynucleotides and Polypeptides of the Invention

FEATURES OF PROTEIN ENCODED BY GENE NO: 1

This gene is expressed primarily in testes tumor and to a lesser extent in fetal brain.

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer particularly of the testes, and defects of the central nervous system such as seizure and neurodegenerative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues of cells, particularly cancer of the testes and central nervous system,

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expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testes and other reproductive tissue, brain and other tissue of the nervous system, and blood cells, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of testicular cancer and treatment of central nervous system disorders since this gene is primarily expressed in the testes tumor and developing brain.

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or bodily fluid from an individual not having the disorder.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 2

This gene is expressed primarily in cancer tissues, such as breast cancer and Wilm's tumor, and to a lesser extent in fetal tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, and/or tumors, particularly, those found in the breast, and developmental

- abnormalities or disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the glandular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., mammary tissue, and fetal tissue and, cancerous and wounded tissues) or bodily fluids (e.g.,
 - tissue, and fetal tissue and, cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 314 as residues: Pro-11 to Thr-18, Leu-43 to Pro-50,
- Gly-64 to Leu-72, and Leu-81 to Lys-86.

 The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of cancers and/or tumors, particularly, those found in the breast since expression is mainly in cancer/tumor
 - corresponding to this gene are useful for treatment/diagnosis of cancers and/or tumor particularly, those found in the breast since expression is mainly in cancer/tumor tissues. May serve as therapeutic proteins for proliferation/differentiation of fetal tissues.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 3

extent in spleen, chronic lymphocytic leukemia. This gene is expressed primarily in CD34 depleted buffy coat and to a lesser

- 7 5 S significantly higher or lower levels may be routinely detected in certain tissues and cell differential identification of the tissue(s) or cell type(s). For a number of disorders of biological sample and for diagnosis of diseases and conditions: blood disorders or cell sample taken from an individual having such a disorder, relative to the standard the above tissues or cells, particularly of the immune system, expression of this gene at directed to these polypeptides are useful in providing immunological probes for leukemias, diseases of the immune system. Similarly, polypeptides and antibodies reagents for differential identification of the tissue(s) or cell type(s) present in a gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an types (e.g., blood cells, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or Therefore, polynucleotides and polypeptides of the invention are useful as
- corresponding to this gene are useful for treatment/diagnosis of blood disorders or leukemias, diseases of the immune system since expression is in tissues related to The tissue distribution indicates that polynucleotides and polypeptides

individual not having the disorder.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 4

This gene is expressed primarily in CD34 depleted buffy coal

ઝ 25 ઝ of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells reagents for differential identification of the tissue(s) or cell type(s) present in a expression level in healthy tissue or bodily sluid from an individual not having the having such a disorder, relative to the standard gene expression level, i.e., the synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, lower levels may be routinely detected in certain tissues and cell types (e.g., blood particularly of the immune system, expression of this gene at significantly higher or polypeptides are useful in providing immunological probes for differential identification lymphocytic diseases. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: blood disorders or Therefore, polynucleotides and polypeptides of the invention are useful as

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expression is in tissues related to immune function. corresponding to this gene are useful for treatment/diagnosis of blood disorders since The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 5

This gene is expressed primarily in CD34 depleted buffy coat

20 5 5 and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or or cell type(s). For a number of disorders of the above tissues or cells, particularly of diseases. Similarly, polypeptides and antibodies directed to these polypeptides are Pro-13 to Lys-21. epitopes include those comprising a sequence shown in SEQ ID NO. 317 as residues spinal fluid) or another tissue or cell sample taken from an individual having such a be routinely detected in certain tissues and cell types (e.g., blood cells, and cancerous the immune system, expression of this gene at significantly higher or lower levels may useful in providing immunological probes for differential identification of the tissue(s) biological sample and for diagnosis of diseases and conditions: blood or immune reagents for differential identification of the tissue(s) or cell type(s) present in a healthy tissue or bodily fluid from an individual not having the disorder. Preferred disorder, relative to the standard gene expression level, i.e., the expression level in Therefore, polynucleotides and polypeptides of the invention are useful as

expression is in tissues related to immune function. corresponding to this gene are useful for treatment/diagnosis of blood disorders since The tissue distribution indicates that polynucleotides and polypeptides

23 FEATURES OF PROTEIN ENCODED BY GENE NO: 6

This gene is expressed primarily in CD34 depleted buffy coat

diseases. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: blood or immune reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

- ၓ or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) be routinely detected in certain tissues and cell types (e.g., and blood cells, and the immune system, expression of this gene at significantly higher or lower levels may
- ઝ cancerous and wounded tissues) or bodily sluids (e.g., serum, plasma, urine, synovial such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 318 as residues: Lys-31 to Lys-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of blood diseases since it is expressed in tissues related to immune function.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 7

This gene is expressed primarily in CD34 depleted buffy coat and to a lesser extent in pineal gland.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases of the immune system and brain associated diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and pineal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another

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fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of blood disorders, immune diseases or brain associated diseases (specifically of the pineal gland) since expression is in tissues related to immune function.

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tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

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FEATURES OF PROTEIN ENCODED BY GENE NO: 8

The translation product of this gene shares sequence homology with an organic cation transporter which is thought to be important in organic cation uptake in the kidney and liver. (See Accession No. 2343059.) Preferred polypeptide fragments comprise the amino acid sequence ITIAIQMICLYNXELYPTFVRNXGVMVCSSLCDIGGITP FIVFRLREVWQALPLIFAVLGLLAAGVTLLLPETKGVALPETMKDAENLGRKAKPKENTTYLK 35 VQTSEPSGT (SEQ ID NO: 615) or TMKDAENLGRKAKPKENT (SEQ ID NO: 616) as well as N-terminal and C-terminal deletions of these fragments. Also preferred are polynucleotide fragments encoding these polypeptide fragments.

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This gene is expressed primarily in liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hepatic and renal diseases where drug elimination/cation exchange (organic cation uptake) in the liver and kidney are problematic. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic or renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., kidney and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression [evel, i.e.,

NO. 320 as residues: Asn-64 to Asn-74, and Gln-81 to Gly-87.

The tissue distribution and homology to organic cation transporter indicate that polynucleotides and polypeptides corresponding to this gene are useful as a polyspecific transporter that is important for drug elimination in the liver (and possibly kidney) since expression is found in the liver.

the expression level in healthy tissue or bodily fluid from an individual not having the

disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID

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FEATURES OF PROTEIN ENCODED BY GENE NO: 9

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This gene is expressed primarily in eosinophil induced with IL-5 and to a lesser extent in fetal liver and spleen. This gene also maps to chromosome 15, and therefore can be used in linkage analysis as a marker for chromosome 15.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases of the immune system, particularly allergies or asthma. Similarly, polypeptides and antibodies directed

to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, liver, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the

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standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating/diagnosis of diseases involving esosinphil reactions since expression seems to be concentrated in eosinophils and other tissues involved in immunity such as the liver and spleen.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 10

This gene is expressed primarily in tissues of hematopoietic lineage and to a lesser extent in Hodgkins lymphoma. Any frame shifts in this sequence can easily be clarified using known molecular biology techniques.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, and immune deficiency or dysfunction. Similarly, polypeptides and

- antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, lymphoid and reticuloendothelial tissues, and cancerous tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/ diagnosis for lymphomas or immune dysfuction or as a therapeutic protein useful in immune modulation based on expression in anergic T-cells and lymphomas.

30 FEATURES OF PROTEIN ENCODED BY GENE NO: 11

This gene is expressed primarily in neutrophils and to a lesser extent in activated lymphoid cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the cell type present in a biological sample and

for diagnosis of diseases and conditions: inflamation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders

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of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 323 as residues: Glu-40 to Lys-46.

10 corresponding to this gene are useful for modulation of an immune reaction or as a growth factor for the differentiation or proliferation of neutrphils for the treatment of neutropenia.

The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 12

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This gene is expressed primarily in brain and to a lesser extent in activated T-cells. It is likely that the open reading frame containing the predicted signal peptide continues in the 5' direction. Preferred polypeptide fragments comprise the amino acid sequence PRVRNSPEDLGLSLTGDSCKL (SEQ ID NO:617).

23 8 30 number of disorders of the above tissues or cells, particularly of the central nervous reagents for differential identification of the tissue(s) or cell type(s) present in a an individual having such a disorder, relative to the standard gene expression level, i.e. plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from detected in certain tissues and cell types (e.g., blood cells, and brain, and other tissue of system, expression of this gene at significantly higher or lower levels may be routinely polypeptides and antibodies directed to these polypeptides are useful in providing disorders including ischemic shock, alzheimers and cognitive disorders. Similarly Val-96, Lys-136 to Ser-145, Ile-152 to Met-169, and Arg-189 to Lys-196 NO. 324 as residues: Ser-5 to Glu-14, Ile-21 to Pro-35, Ser-65 to Asp-81, Cys-89 to disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID the expression level in healthy tissue or bodily fluid from an individual not having the the nervous system and cancerous and wounded tissues) or bodily fluids (e.g., serum, immunological probes for differential identification of the tissue(s) or cell type(s). For a biological sample and for diagnosis of diseases and conditions: neurodegenerative Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnostic/treatment for cancers of the given tissue or in the treatment of neurological disorders of the CNS.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 13

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This gene was also recently cloned by other groups, naming this calcium-activated potassium channel gene, hKCa4. (See Accession No. AF033021, see also, Accession No. 2584866.) This gene is mapped to human chromosome 19q13.2. A second signal sequence likely exists upstream from the predicted signal sequence as described in Table 1. Preferred polypeptide fragments comprise: QADDLQATVAALCVLRGGGPWAG SWLSPKTPGAMGGDLVLGLGALRRRKRLL (SEQ NO: 618); or EQEKSLAGWALVLAXXGIGL MVLHAEMLWFGGCSAVNATGHLSDTLWLIPTFLTIGYGDVVPCTMWGKIVCLCTGVMGVCC TALLVAVVARKLEFNKAEKHVHNFMMDIQYTKEMKESAARVLQEAWMFYKHTRRKESHAAR XHQRXLLAANNAFRQVRLKHRKLREQVNSMVDISKMHMILYDLQQNLSSSHRALEKQIDTLAG KLDALTELLSTALGPRQLPEPSQQSK (SEQ ID NO: 619), as well as N-terminal and C-terminal deletions. Also preferred are polynulcleotide fragments encoding these polypeptide fragments.

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15 This gene is expressed primarily in breast lymph node and T-cells, and to a lesser extent in placenta.

immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides lower levels may be routinely detected in certain tissues and cell types (e.g., lymphoid tissue, blood cells and placenta, and cancerous and wounded tissues) or bodily fluids sample taken from an individual having such a disorder, relative to the standard gene particularly of the immune system, expression of this gene at significantly higher or expression level, i.e., the expression level in healthy tissue or bodily fluid from an (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Therefore, polynucleotides and polypeptides of the invention are useful as ndividual not having the disorder. Preferred epitopes include those comprising a are useful in providing immunological probes for differential identification of the biological sample and for diagnosis of diseases and conditions: hematologic and issue(s) or cell type(s). For a number of disorders of the above tissues or cells, reagents for differential identification of the tissue(s) or cell type(s) present in a sequence shown in SEQ ID NO. 325 as residues: Arg-13 to Lys-23. 25 8 3

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment/diagnosis of hematologic and diseases involving immune modulation based or distribution in the lymph node and T-cells.

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FEATURES OF PROTEIN ÉNCODED BY GENE NO: 14

This gene was recently cloned by another group, calling it PAPS synethase. (See Accession No. e1204135.) Preferred polypeptide fragments comprise the amino acid sequence YQAHHVSRNKRGQVVGTRGGFRGCTVWLTGLSGAGK (SEQ ID NO: 620).

Also preferred are the polynucleotide fragments encoding this polypeptide fragment.

It has been discovered that this gene is expressed primarily in benign prostate hyperplasia, Human Umbilical Vein Endothelial Cells and to a lesser extent in smooth muscle and Human endometrial stromal cells-treated with estradiol.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: inflamation, ischemia, and restenosis, based on endothelial cell and smooth muscle cell expression, and prostate diseases such as benign prostate hyperplasia or prostate cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate or vessels of the circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., prostate, endothelial cells, smooth muscle, and endometrium, and cancerous and wounded tissues) or bodily

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20 fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 326 as residues: Arg-21 to Asp-26, Lys-35 to Lys-44, 25 Glu-49 to Asn-58.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating/diagnosing diseases or conditions where the endothelial cell lining of the veins and arteries of underlying smooth muscle are involved.

FEATURES OF PROTEIN ENCODED BY GENE NO: 15

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This gene is expressed primarily in human 6 week embryo and to a lesser extent in placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: developmental anomalies or fetal deficiencies. Similarly, polypeptides and antibodies directed to these

an individual having such a disorder, relative to the standard gene expression level, i.e., disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from tissue, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, the expression level in healthy tissue or bodily fluid from an individual not having the lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic particularly developmental in nature, expression of this gene at significantly higher or polypeptides are useful in providing immunological probes for differential identification

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corresponding to this gene are useful for detection of developmental abnormalities The tissue distribution indicates that polynucleotides and polypeptides 5

NO. 327 as residues Lys-50 to Glu-57.

FEATURES OF PROTEIN ENCODED BY GENE NO: 16

2 linkage analysis as a marker for chromosome 14. in fetal tissues. This gene is mapped to chromosome 14, and therefore is useful in This gene is expressed primarily in kidney and amygdala and to a lesser extent

system or developing fetal tissues, expression of this gene at significantly higher or of the tissue(s). For a number of disorders of the above tissues, particularly of the renal polypeptides are useful in providing immunological probes for differential identification developmental abnormalities. Similarly, polypeptides and antibodies directed to these for diagnosis of diseases and conditions: kidney diseases, neurological disorders and reagents for differential identification of the tissue(s) present in a biological sample and Therefore, polynucleotides and polypeptides of the invention are useful as

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serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample arnygdala, and fetal tissues, and cancerous and wounded tissues) or bodily fluids (e.g., lower levels may be routinely detected in certain tissues and cell types (e.g., kidney, individual not having the disorder. expression level, i.e., the expression level in healthy tissue or bodily fluid from an taken from an individual having such a disorder, relative to the standard gene

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corresponding to this gene are useful for treatment or diagnosis of conditions affecting the brain, kidneys and fetal development The tissue distribution indicates that polynucleotides and polypeptides 30

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This gene is expressed primarily in ovarian cancer

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5 plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from the reproductive system. expression of this gene at significantly higher or lower levels or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) ovarian cancer Similarly, polypeptides and antibodies directed to these polypeptides are the expression level in healthy tissue or bodily fluid from an individual not having the an individual having such a disorder, relative to the standard gene expression level, i.e. reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, may be routinely detected in certain tissues and cell types (e.g., ovarian and other biological sample and for diagnosis of diseases and conditions; solid tumors similar to reagents for differential identification of the tissue(s) or cell type(s) present in a NO. 329 as residues Ser-51 to Val-56. disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID Therefore, polynucleotides and polypeptides of the invention are useful as

corresponding to this gene are useful for the treatment of solid tumors of the reproductive system such as ovarian cancer. The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 18

GLTLTTCSGPTEKPATKNYFLKRLLQEMHIRAN (SEQ ID NO: 644), as well as N-terminal polypeptide fragments. and C-terminal deletions. Also preferred are polynucleotide fragments encoding these ILILPYCAHLHEELNC (SEQ ID NO: 643) and SFFISEEKGHLLLQAERHPWYAGALYGYSG fragments comprise the amino acid sequence: IRHEQHPNFSLEMHSKGSSLLLFLPQL This gene is expressed primarily in brain medulloblastoma. Preferred polypeptide

25 20 30 or cell type(s). For a number of disorders of the above tissues or cells, particularly of biological sample and for diagnosis of diseases and conditions: tumors particularly of sample taken from an individual having such a disorder, relative to the standard gene tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids levels may be routinely detected in certain tissues and cell types (e.g., brain and other the Central nervous system, expression of this gene at significantly higher or lower useful in providing immunological probes for differential identification of the tissue(s) the CNS or Similarly, polypeptides and antibodies directed to these polypeptides are reagents for differential identification of the tissue(s) or cell type(s) present in a (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Therefore, polynucleotides and polypeptides of the invention are useful as

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating medulloblastoma or similar tumors.

FEATURES OF PROTEIN ENCODED BY GENE NO: 19

This gene is expressed primarily in adipocytes.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: obesity. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the adipose tissues expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adipocytes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating obesity by regulating the function and number of adipocytes

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FEATURES OF PROTEIN ENCODED BY GENE NO: 20

This gene is expressed primarily in B cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, of the immune system with an emphasis on B cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumors of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of B cell derived tumors based on its expression in b cell lymphomas

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FEATURES OF PROTEIN ENCODED BY GENE NO: 21

This gene is expressed primarily in immune cells and to a lesser extent in fetal

tins gene is expressed primarity in minimum

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: inflammatory diseases Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cells of the immune system; and fetal tissues, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an

individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO:333 as residues Asp-10 to Pro-19, Ser-74 to Tyr-79, Glu-95 to Lys-110.

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The tissue distribution indicates that polynucleotides and polypeptides

corresponding to this gene are useful for treatment of diseases involving alterations in T

FEATURES OF PROTEIN ENCODED BY GENE NO: 22

It has been discovered that this gene is expressed primarily in ovarian tumor. Therefore, polynucleotides and polypeptides of the invention are useful as

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors particularly of the ovary. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of tumors of the reproductive organs. expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., ovarian

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and other reproductive tissue and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 334 as residues: Leu-22 to Gln-27.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of ovarian tumors as it has only been identified in ovarian tumors.

FEATURES OF PROTEIN ENCODED BY GENE NO: 23

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It has been discovered that this gene is expressed primarily in fetal tissues and to a lesser extent in osteoclastoma cell line

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: osteoporosis or arthritis Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the

skeletal expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone cells, and fetal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of conditions of abnormal bone remodeling due to enhanced activity of osteoclasts. This may be useful as a specific marker for malignancies derived from osteoclasts or their precursors.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 24

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The translation product of this gene shares sequence homology with a periplasmic ribonuclease which is thought to be important in degrading extracellular polynucleotides

It has been discovered that this gene is expressed primarily in serum treated smooth muscle cells

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5 S spinal fluid) or another tissue or cell sample taken from an individual having such a the vasculature expression of this gene at significantly higher or lower levels may be or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) restenosis. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: vascular disease such as reagents for differential identification of the tissue(s) or cell type(s) present in a healthy tissue or bodily fluid from an individual not having the disorder. Preferred routinely detected in certain tissues and cell types (e.g., smooth muscle, and cancerous Gln-30 to Lys-36, and Pro-41 to Arg-48. epitopes include those comprising a sequence shown in SEQ ID NO: 336 as residues: disorder, relative to the standard gene expression level, i.e., the expression level in and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution and homology to ribonucleases indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of pathological conditions of smooth muscle associated with bacterial or viral infiltration

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FEATURES OF PROTEIN ENCODED BY GENE NO: 25

20 မွ 25 cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial expression of this gene at significantly higher or lower levels may be routinely detected tissues or cells, particularly of the human brain development and related diseases, identification of the tissue(s) or cell type(s). For a number of disorders of the above these polypeptides are useful in providing immunological probes for differential development and related diseases. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: human brain reagents for differential identification of the tissue(s) or cell type(s) present in a such a disorder, relative to the standard gene expression level, i.e., the expression level in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and in healthy tissue or bodily fluid from an individual not having the disorder. fluid or spinal fluid) or another tissue or cell sample taken from an individual having This gene is expressed primarily in Early Stage Human Brain Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution and homology to this gene indicate that polynucleotides
and polypeptides corresponding to this gene are useful for diagnosis and treatment of
diseases affecting human brain development and related diseases.

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It has been discovered that this gene is expressed primarily in human brain tissue

FEATURES OF PROTEIN ENCODED BY GENE NO: 26

biological sample and for diagnosis of diseases and conditions: human brain diseases and other diseases related to brain diseases, which may be caused by brain diseases. Therefore, polynucleotides and polypeptides of the invention are useful as Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a

may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from the expression level in healthy tissue or bodily fluid from an individual not having the human brain diseases, expression of this gene at significantly higher or lower levels providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the 2 2

The tissue distribution and homology to the gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of human brain diseases and other diseases related. 20

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FEATURES OF PROTEIN ENCODED BY GENE NO: 27

It has been discovered that this gene is expressed primarily in Anergic T-cells.

inflammatory diseases and diseases related to T lymph cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell inflammatory diseases and diseases related to T lymph cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: immune diseases, reagents for differential identification of the tissue(s) or cell type(s) present in a disorders of the above tissues or cells, particularly of the immune diseases, 30 23

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. The tissue distribution and homology to the gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for immune diseases,

inflammatory diseases and diseases related to Tlymph cells. S

FEATURES OF PROTEIN ENCODED BY GENE NO: 28

The translation product of this gene shares sequence homology with Shigella flexneri positive transcriptional regulator CriR (criR) gene which is thought to be important in regulation of gene expression.

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This gene is expressed primarily in human synovial sarcoma and normal human brain tissues.

particularly sarcomas of the synovium. Similarly, polypeptides and antibodies directed issues or cells, particularly of the human brain and synovium and other related human biological sample and for diagnosis of diseases and conditions: human brain diseases identification of the tissue(s) or cell type(s). For a number of disorders of the above these polypeptides are useful in providing immunological probes for differential Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 2

an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from brain diseases, expression of this gene at significantly higher or lower levels may be nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, routinely detected in certain (e.g., synovial tissue, and brain and other tissue of the

the expression level in healthy tissue or bodily fluid from an individual not having the 23

corresponding to this gene are useful for diagnosis and treatment of human synovial The tissue distribution indicates that polynucleotides and polypeptides sarcoma and other related human brain diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 29

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prostate and to a lesser extent in pineal gland, adipose tissue, kidney, adrenal gland, This gene is expressed in bone marrow, infant brain, fetal liver and spleen, umbilical vein endothelial cells, and T cells.

reagents for identification of the tissue(s) or cell type(s) present in a biological sample Therefore, polynucleotides and polypeptides of the invention are useful as and for diagnosis of diseases and conditions: diseases related to bone marrow or

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types (e.g., blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample

taken from an individual having such a disorder, relative to the standard gene

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hematoplastic tissues, prostate, kidney, adrenal gland, and cardiovascular tissue or organs. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the diseases related to

- 5 hematoplastic tissues, immune system, prostate, kidney, adrenal gland, and cardiovascular tissue or organs, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, hematopoietic cells, pineal gland, adipose tissue, kidney, adrenal gland, endothelial cells, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene
- The tissue distribution and homology to the gene indicate that polynucleotides

 15 and polypeptides corresponding to this gene are useful for diagnosis and treatment of

 diseases related to hematoplastic tissues, immune system, prostate, kidney, adrenal

 gland, and cardiovascular tissue or organs.

expression level, i.e., the expression level in healthy tissue or bodily fluid from an

individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 30

20 This gene is expressed primarily in meningea and to a lesser extent in breast and adult brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Diseases of the meanings and related brain diseases. Similarly polypeptides and antibodies directed

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meningea and related brain diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the meningea and related brain diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., miningea, mammary tissue, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., scrum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the meningea and related brain diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 3

This gene is expressed in meningea, fetal spleen, osteoblast and to a lesser extent in activated T-cells, endometrial stromal cells, fetal lung, HL-60, thymus, testis and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: meningeal disease, osteoporosis, immune diseases, and hematoplastic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the

diseases, and hematoplastic diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, endometrium, lung, thymus, testis, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of meningeal, osteoporosis, immune diseases, hematoplastic diseases, testis diseases and

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lung diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 32

This gene is expressed primarily in human thymus and to a much lesser extent in infant brain, T-cells, smooth muscle, endothelial cells, bone marrow, human ovarian tumor and keratinocytes testes, osteoclastoma, breast, and tonsils.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Diseases involving the thymus, particularly thymic cancer and diseases involving T-cell maturation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing

immunological probes for differential identification of the tissue(s) or cell type(s). For a

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number of disorders of the above tissues or cells, particularly of the thymus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., thymus, brain, and other tissue of the nervous system, blood cells, bone marrow, ovaries, and testes, and other reproductive tissue, mammary tissue, tonsils, melanocytes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the thymus particularly thymic cancer and diseases involving T-cell maturation.

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15 FEATURES OF PROTEIN ENCODED BY GENE NO: 33

This gene is expressed primarily in human tonsils, and placenta, and to a lesser extent in adipocytes, melanocyte, and infant brain.

polypeptides are useful in providing immunological probes for differential identification gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an particularly of the inflammatory diseases, immune diseases, and obesity, expression of biological sample and for diagnosis of diseases and conditions: inflammatory diseases, immune diseases, and obesity. Similarly, polypeptides and antibodies directed to these of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, tissues and cell types (e.g., tonsils, placenta, adipocytes, melanocytes, and brain and fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard this gene at significantly higher or lower levels may be routinely detected in certain other tissue of the nervous system, and cancerous and wounded tissues) or bodily Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. ಣ 25 3

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The tissue distribution and homology to this gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases such as inflammation, immune diseases, and obesity.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 34

This gene is expressed in activated T cells, and to a lesser extent in pituitary, testis, and breast lymph node.

disorders of the immune system, expression of this gene at significantly higher or lower cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in levels may be routinely detected in certain tissues and cell types (e.g., pituitary, testes biological sample and for diagnosis of diseases and conditions: diseases relating to T and other reproductive tissue, mammary tissue, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or providing immunological probes for differential identification of the tissue(s) or cell pinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in ype(s). For a number of disorders of the above tissues or cells, particularly, of the Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a nealthy tissue or bodily fluid from an individual not having the disorder. S 2 15

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of immune disorders.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 35

This gene is expressed primarily in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the diseases relating to neurological disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and celltypes (e.g., brain, and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological

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This gene is expressed primarily in infant brain

FEATURES OF PROTEIN ENCODED BY GENE NO: 36

S providing immunological probes for differential identification of the tissue(s) or cel Similarly, polypeptides and antibodies directed to these polypeptides are useful in biological sample and for diagnosis of diseases and conditions: neurological disorders. reagents for differential identification of the tissue(s) or cell type(s) present in a type(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful

2 5 standard gene expression level, i.e., the expression level in healthy tissue or bodily diseases relating to neurological disorders, expression of this gene at significantly bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another brain and other tissue of the nervous system, and cancerous and wounded tissues) or higher or lower levels may be routinely detected in certain tissues and cell types (e.g., tissue or cell sample taken from an individual having such a disorder, relative to the

corresponding to this gene are useful for diagnosis and treatment of neurological The tissue distribution indicates that polynucleotides and polypeptides fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 37

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This gene is expressed primarily in human ovary.

providing immunological probes for differential identification of the tissue(s) or cel biological sample and for diagnosis of diseases and conditions: ovarian cancer. type(s). For a number of disorders of the above tissues or cells, particularly of the Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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ઝ expression of this gene at significantly higher or lower levels may be routinely detected in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial in certain tissues and cell types (e.g., ovary and other reproductive tissue, and ovarian disorders such as those involving germ cells, ovarian follicles, stromal cells,

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corresponding to this gene are useful for diagnosis and treatment of ovariopathy The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 38

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in lymph node breast cancer.

5 expression of this gene at significantly higher or lower levels may be routinely detected in healthy tissue or bodily fluid from an individual not having the disorder. polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: breast cancer. Similarly, such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily sluids (e.g., serum, plasma, urine, synovial in certain tissues and cell types (e.g., mammary tissue and lymphoid tissue, and number of disorders of the above tissues or cells, particularly of the breast cancer, immunological probes for differential identification of the tissue(s) or cell type(s). For a

corresponding to this gene are useful for used as a diagnostic marker for breast cancer. The tissue distribution indicates that polynucleotides and polypeptides 5

FEATURES OF PROTEIN ENCODED BY GENE NO: 39

25 20 30 biological sample and for diagnosis of diseases and conditions; neuronal disorders such the above tissues or cells, particularly of the brain, expression of this gene at differential identification of the tissue(s) or cell type(s). For a number of disorders of directed to these polypeptides are useful in providing immunological probes for as trauma, brain degeneration, and brain tumor. Similarly, polypeptides and antibodies reagents for differential identification of the tissue(s) or cell type(s) present in a another tissue or cell sample taken from an individual having such a disorder, relative to tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or significantly higher or lower levels may be routinely detected in certain tissues and cell the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. types (e.g., brain and other tissue of the nervous system, and cancerous and wounded Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in brain and to a lesser extent in other tissues.

႘ corresponding to this gene are useful for diagnosis and therapeutic treatment of The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 40

This gene is expressed in early stage human embryo, adrenal gland tumor, and immune tissues such as fetal liver, fetal spleen, T-cell, and myoloid progenitor cell line and to a lesser extent in ovary, colon cancer, and a few orther tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumorigenesis including adrenal gland tumor, colon cancer and various other tumors, developmental and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer tissues, early stage human tissues, and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, blood cells, bone marrow, ovary and other reproductive tissue, and colon, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and therapeutic treatment of immune and developmental disorders, and tumorigenesis.

gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

individual not having the disorder.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 41

This gene is expressed primarily in fetal lung, endothelial cells, liver, thymus and a few other immune tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as

oreagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune disorders such as immune deficiency and autoimmune diseases, pulmonary diseases, liver diseases, and tumor matasis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal lung, liver, endothelial cells, and immune tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain

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tissues and cell types (e.g., lung, endothelial cells, liver, thymus, and other tissue of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of immune disorders and pulmonary and hepatic diseases. Its promoter may also be used for immune system and lung-specific gene therapies. The expression of this gene in endothelial cells indicates that it may also involve in angiogenesis which therefore may play role in tumor matasis.

FEATURES OF PROTEIN ENCODED BY GENE NO: 42

This gene is expressed primarily in liver, thyroid, parathyroid and to a lesser extent in fetal lung, stomach and early embryos.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: metabolic regulation, obesity, heptic failure, heptacellular tumors or thyroiditis and thyroid tumors. Similarly, polymentides and antihodies directed to these polymentides are useful in providing

- polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive/endocrine system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, thyroid, parathyroid, lung, stomach, and embryonic tissue, and cancerous and wounded tissues) or bodily fluids
 - stomach, and embryonic tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- 30 The tissue distribution and the extracellular locations indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of digestive/endocrine disorders, including metabolic regulation, heptic failure, malabsortion, gastritis and neoplasms.

FEATURES OF PROTEIN ENCODED BY GENE NO: 43

cortex, hypothalmus and to a lesser extent in retina, adipose and stomach cancer and This gene is expressed primarily in Schizophrenic adult brain, pituitary, front

- 2 5 S of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells cell sample taken from an individual having such a disorder, relative to the standard or lower levels may be routinely detected in certain tissues and cell types (e.g., retinal particularly of the central nerve system, expression of this gene at significantly higher biological sample and for diagnosis of diseases and conditions: schizophrenia and other reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. polypeptides are useful in providing immunological probes for differential identification neurological disorders. Similarly, polypeptides and antibodies directed to these gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an tissue, adipose, stomach, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or Therefore, polynucleotides and polypeptides of the invention are useful as
- system, including schizophrenia, neurodegeneration, and neoplasia. Additionally, a corresponding to this gene are useful in treatment/detection of disorders in the nerve secreted protein in brain may serve as an endocrine. The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 44

binding proteins which are thought to be important in signal transduction and protein transport. The translation product of this gene shares sequence homology with GTP

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and to a lesser extent in gall bladder. cells, GM-CSF treated macrophage, anergic T cells, osteoblast, osteoclast, CD34+ cells This gene is expressed primarily in umbilical vein and microvascular endothelial

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immunological probes for differential identification of the tissue(s) or cell type(s). For a growth, osteonecrosis, osteoporosis, angiogenesis and/or hematopoeisis. Similarly, biological sample and for diagnosis of diseases and conditions: bone formation and reagents for differential identification of the tissue(s) or cell type(s) present in a hematopoeisis systems, expression of this gene at significantly higher or lower levels number of disorders of the above tissues or cells, particularly of the skeletal and polypeptides and antibodies directed to these polypeptides are useful in providing Therefore, polynucleotides and polypeptides of the invention are useful as

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an sample taken from an individual having such a disorder, relative to the standard gene (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell cells, bone, and gall bladder, and cancerous and wounded tissues) or bodily fluids may be routinely detected in certain tissues and cell types (e.g., endothelial cells, blood individual not having the disorder.

hematopoeisis because its involvement in the growth signaling or angiogenesis. polynucleotides and polypeptides corresponding to this gene are useful for treatment/detection of bone formation and growth, osteonecrosis, osteoporosis, and/or The tissue distribution and homology to GTP binding proteins indicate that

FEATURES OF PROTEIN ENCODED BY GENE NO: 45

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sequence receptor gamma subunit which is thought to be important in protein The translation product of this gene shares sequence homology with signal

5 translocation on endoplasmic reticulum.

to a lesser extent in endothelial cells and smooth muscle. This gene is expressed primarily in adrenal gland, salivary gland, prostate, and

- 20 providing immunological probes for differential identification of the tissue(s) or cell biological sample and for diagnosis of diseases and conditions: protein secretion. secretory organs, expression of this gene at significantly higher or lower levels may be Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a type(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful as
- 25 ઝ routinely detected in certain tissues and cell types (e.g., adrenal gland, salivary gland, fluid from an individual not having the disorder prostate, endothelial cells, and smooth muscle, and cancerous and wounded tissues) or standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another

disorders, prostate cancer, xerostomia or sialorrhea polynucleotides and polypeptides corresponding to this gene are useful for endocrine The tissue distribution and homology to SSR gamma subunit indicate that

ઝ FEATURES OF PROTEIN ENCODED BY GENE NO: 46

melanocyte, amygdala, brain, and stomach. This gene is expressed primarily in osteoclastoma cells and to a lesser extent in

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: ossification, osteoporosis, fracture, osteonecrosis, osteosarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., melanocytes, amygdala, brain and other tissue of the nervous system,

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and stomach, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful in intervention of ossification, osteoporosis, fracture, osteonecrosis and osteosarcoma.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 48

20 The translation product of this gene shares sequence homology with proline rich proteins which is thought to be important in protein-protein interaction.

This gene is expressed primarily in brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological and psychological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nerve system and endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to proline-rich proteins indicate that polynocleotides and polypeptides corresponding to this gene are useful in intervention

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and detection of neurological diseases, including trauma, neoplasia, degenerative or metabolic conditions in the central nerve system. Additionally, the gene product may be a secreted by the brain as an endocrine.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 49

The translation product of this gene shares sequence homology with the AOCB gene from Aspergillus nidulans which is important in asexual development.

This gene is expressed primarily in infant brain and to a lesser extent in the developing embryo, trachea tumors, B-cell lymphoma and synovial sarcoma.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurodegenerative diseases, leukemia and sarcoma's. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, blood cells, trachea, and synovial tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or
- healthy tissue or bodily fluid from an individual not having the disorder.

 The tissue distribution in infant brain and sarcoma's and homology to a gene involved in a key step of eukaryotive development (fungal spore formation) indicates that the protein product of this clone could play a role in neurological diseases such as

spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in

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involved in a key step of eukaryotivc development (fungal spore formation) indicates that the protein product of this clone could play a role in neurological diseases such as schizophrenia, particularly in infants. The existence of the gene in a B-cell lymphoma indicates the gene may be used in the treatment and detection of leukernia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 50

30 This gene is expressed primarily in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: pulmonary disorders including lung cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the pulmonary system, expression of this gene at significantly higher or

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lower levels may be routinely detected in certain tissues and cell types (e.g., lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution of this gene only in fetal lung indicates that it plays a key role in development of the pulmonary system. This would suggest that misregulation of the expression of this protein product in the adult could lead to lymphoma or sarcoma formation, particularly in the lung. It may also be involved in predisposition to certain pulmonary defects such as pulmonary edema and embolism, bronchitis and cystic fibrosis.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 51

This gene is expressed primarily in hematopoietic cell types and fetal cells and to

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a lesser extent in all tissue types.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions; defects in the immune system and hematopoeisis. Similarly, polypeptides and antibodies directed to these

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hematopoietic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, and fetal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

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The tissue distribution of this gene predominantly in hematopoeitic cells and in the developing embryo indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection and treatment of lymphomas and disease states affecting the immune system or hematopoeisis disorders such as leukemia, AIDS, arthritis and asthma.

fluid from an individual not having the disorder.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 52

This gene is expressed primarily in prostate and to a lesser extent in fetal spleen, fetal liver, infant brain and T cell leukemias.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: prostate disorders, prostate cancer, leukemia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, and/or prostate gland expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., thymus, spleen, liver, brain and other tissue of the nervous system, and

blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution of this gene in prostate indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection or treatment of prostate disorders or prostate cancer. Its distribution in fetal liver and fetal spleen indicates it may play a role in the immune system and its misregulation could lead to immune disorders such as leukemia, arthritis and asthma.

FEATURES OF PROTEIN ENCODED BY GENE NO: 53

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The translation product of this gene shares sequence homology with dynein. This gene is expressed primarily in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosts of diseases and conditions: neuro-degenerative diseases of the brain. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

- or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an
- 35 gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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molecules indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection/treatment of neurodegenerative diseases, such as Alzheimers, The predominant tissue distribution in the brain and homology to dynein, a microtubule motor protein involved in the positioning of cellular organelles and Huntigtons, Parkinsons diseases and shizophrenia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 54

The translation product of this gene shares sequence homology with ubiquitinconjugation protein, an enzyme which is thought to be important in the processing of the Huntingtons Disease causing gene.

This gene is expressed primarily in brain and to a lesser extent in activated

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial significantly higher or lower levels may be routinely detected in certain tissues and cell disease states including Huntington's disease. Similarly, polypeptides and antibodies biological sample and for diagnosis of diseases and conditions: neurodegenerative differential identification of brain tissues. For a number of disorders of the above lissues or cells, particularly of the neurological systems expression of this gene at directed to these polypeptides are useful in providing immunological probes for types (e.g., brain and other tissue of the nervous system, and blood cells, and 2 8

such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having in healthy tissue or bodily fluid from an individual not having the disorder. 25

The predominant tissue distribution of this gene in the brain and its homology to muscular atrophy. In addition, the existence of elevated levels of free ubiquitin pools in Alzheimer's disease, Parkinson's disease and amylotrophic lateral sclerosis indicates Thus, considering the gene described here is homologous to a ubiquitin-conjugation that the ubiquitin pathway of protein degradation plays a role in these disease states. spinocerebullar ataxia types I and III, dentatorubropallidoluysian and spinal bulbar a Huntington interacting protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the regulation of the expression of the Huntington disease gene and other neurodegenerative diseases including protein it may play a general role in neurodegenarative conditions.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 56

This gene is expressed primarily in T-cells (anergic T-cells, resting|T-Cells, apoptotic T-cells) and lymph node (breast), as well as brain (hypothalamus hippocampus, pituitary, infant brain, early-stage brain).

immunological probes for differential identification of the tissue(s) or cell type(s). For a cells, lymphoid tissue, and brain and other tissue of the nervous system, and cancerous neurological (e.g. Alzheimer's disease, dementia, schizophrenia) disorders, Similarly, hematopoietic and immune systems, expression of this gene at significantly higher or number of disorders of the above tissues or cells, particularly of the central nervous, and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in immunodeficiencies, autoimmunities, inflammation, leukemias & lymphomas) and lower levels may be routinely detected in certain tissues and cell types (e.g., blood Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and antibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune (e.g. healthy tissue or bodily fluid from an individual not having the disorder. 2 12 S

associated with the hematopoietic and immune systems, such as anemias (leukemias). developmental brain defects, neuro-degenerative diseases or behavioral abnomalities corresponding to this gene are useful in the intervention or detection of pathologies The tissue distribution indicates that polynucleotides and polypeptides In addition, the expression in brain (including fetal) might suggest a role in (e.g. schizophrenia, Alzheimer's, dementia, depression, etc.). 8

FEATURES OF PROTEIN ENCODED BY GENE NO: 57

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This gene is expressed primarily in lung, and to a lesser extent in a variety of other hematological cell types (e.g. Raji cells, bone marrow cell line, activated monocytes).

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polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vasculo-pulmonary and hematopoietic systems, expression of this hematological disfunction. Similarly, polypeptides and antibodies directed to these Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: pulmonary and/or reagents for differential identification of the tissue(s) or cell type(s) present in a

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gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., lung and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful in the intervention and detection of pathologies associated with the vasculo-pulmonary system. In addition the expression of this gene in a variety of leukocytic cell types and a bone marrow cell line might suggest a role in hematopoietic and immune system disorders, such as leukemias & lymphomas, inflammation, immunodeficiencies and autoimmunities.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 58

The translation product of this gene shares sequence homology with adenylate kinase isozyme 3 (gill 63528 GTP:AMP phosphotransferase (EC 2.7.4.10) [Bos taurus]), which is thought to be important in catalyzing the phosphorylation of AMP to ADP in the presence of ATP or inorganic triphosphate.

This gene is expressed primarily in fetal liver, heart and placenta, and to a lesser extent in many other tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hepatic, cardiovascular or reproductive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic, cardiovascular and reproductive systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, heart, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of conditions related to hepatic function and pathogenesis, in particular, those dealing with liver development and the differentiation of hepatocyte progenitor cells.

from an individual not having the disorder.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 59

This gene is expressed primarily in CD34 positive cells (Cord Blood).

Therefore, polynucleotides and polypeptides of the invention are useful as

Therefore, polynucleotides and polypeptides of the invention are useful as

- biological sample and for diagnosis of diseases and conditions: hematopoietic differentiation and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily
- fluid from an individual not having the disorder.

 The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful in the detection and treatment of conditions associated with CD34-positive cells, and therefore as a marker for cell differentiation in

20 hematapoiesis, as well as immunological disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 60

The translation product of the predicted open reading frame of this contig has sequence identity to the murine gene designated Insulin-Like Growth Factor-Binding Protein (IGFBP)-1 as described by Lee and colleagues (Hepatology 19 (3), 656-665

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This gene is expressed exclusively in hemangiopericytoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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- biological sample and for diagnosis of hemangiopericytoma and other pericyte or endothelial cell proliferative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the circulatory and immune systems, expression of this tissues or cells, particularly of the circulatory and immune systems.
- 35 gene at significantly higher or lower levels may routinely be detected in certain tissues and cell types (e.g., pericyte or endothelial cells, and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

Polynucleotides and polypeptides corresponding to this gene are useful as cell growth regulators since IGFBP-1-like molecules function as modulators of insulin-like growth factor activity. In addition, since IGFBP-1 is expressed at high levels following hepatectomy and during fetal liver development, polynucleotides of the present invention may also be used for the diagnosis of developmental disorders. Further, polypeptides of the present invention may be used therapeutically to treat developmental liver disorders as well as to regulate hepatocyte and supporting cell growth following hepatectomy or to treat liver disorders.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hemangiopericytoma and liver disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 61

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This gene is expressed primarily in schizophrenic frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: nervous system and cognitive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the frontal cortex and CNS expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, treatment and diagnosis of frontal cortex, neuro-degenerative and CNS disorders

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35 FEATURES OF PROTEIN ENCODED BY GENE NO: 62

This gene is expressed primarily in human adrenal gland tumor, and to a lesser extent in human kidney, medulla and adult pulmonary tissue.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: metabolic endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are

- or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine and nervous system disorders and neoplasia, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adrenal gland, kidney, brain and other tissue of the nervous system, nulmonary tissue and cancerous and wounded tissues) or hodily fluids (e.g., serum
- 10 pulmonary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, treatment and diagnosis of neurological and endocrine disorders including neoplasia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 63

20 This gene is expressed primarily in human adipocytes, and to a lesser extent in spleen, 12-week old human, and testes.

spleen, 12-week old human, and testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune, metabolic and growth disorders. Similarly, polypeptides and antibodies directed to these polypeptides

- are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types(e.g., adipocytes, spleen, and testes and other reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that polynucleotides and polypeplides corresponding to this gene are useful for study, diagnosis and treatment of immune, developmental and metabolic disorders.

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PZF. (See Accession No. 453376; see also Gene 152 (2), 233-238 (1995).) Preferred One translated product of this clone is homologous to the mouse zinc finger protein FEATURES OF PROTEIN ENCODED BY GENE NO: 64

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SAGP (SEQ ID NO: 624); IQYVRCEMEGCGTVLAHPRYLQHHIKYQHLLKKKYVCPHPSCGRLF mouse and man. For example, preferred polypeptide fragments comprise the amino acid ID NO: 626); RSSRTSVSRHRDTENTRSSRSKTGSLQLICKSEPNTDQLDY (SEQ ID NO: 627); RLQKQLLRHAKHHT (SEQ ID NO: 625); DQRDYICEYCARAFKSSHNLAVHRMIHTGEK (SEQ SKSY (SEQ ID NO: 623); CSGTERVSLMADGKIFVGSGSSGGTEGLYMNSDILGATTEVLIEDSD sequence: LQCEICGFTCRQKASLNWHMKKHDADSFYQFSCNICGKKFEKKDSYVAHKAKSH polypeptide fragments correspond to the highly conserved domains shared between HHIKYQHLLKKKYYCPHPSCGRLFRLQKQLLRHAKHHTD (SEQ ID NO: 629); or residues PFKDDPRDETYKPHLERETPKPRRKSG (SEQ ID NO: 630); QYVRCEMEGCGTVLAHPRYLQ PEV (SEQ ID NO: 621); ITSTDILGTNPESLTQPSD (SEQ ID NO: 622); NSTSGECLLLEAEGM

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5 are polynucleotide fragments encoding these polypeptide fragments. 151-182 of QRDYICEYCARAFKSSHNLAVHRMIHTGEKHY (SEQ ID NO: 628). Also preferred

cancer tissue and to a lesser extent in smooth muscle, pancreatic tumor, and apoptotic This gene is expressed primarily in Rhabdomyosarcoma, melanocyte and colon

- 23 20 or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) biological sample and for diagnosis of diseases and conditions which include, but are not limited to.. Similarly, polypeptides and antibodies directed to these polypeptides are reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as
- levels may be routinely detected in certain tissues and cell types (e.g., striated muscle. the immune and hemopoetic, expression of this gene at significantly higher or lower or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal melanocytes, colon, smooth muscle, pancreas, and blood cells, and cancerous and

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corresponding to this gene are useful for study, diagnosis and treatment of cancer and The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 65

to a lesser extent in human bone marrow and fetal kidney. This gene is expressed primarily in human adipose and salivary gland tissue and

reagents for differential identification of the tissue(s) or cell type(s) present in a useful in providing immunological probes for differential identification of the tissue(s) disorders. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: metabolic and immune Therefore, polynucleotides and polypeptides of the invention are useful as

- 5 or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and hemopoetic systems, expression of this gene at significantly higher or standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another salivary gland, bone marrow, and kidney, and cancerous and wounded tissues) or lower levels may be routinely detected in certain tissues and cell types (e.g., adipose,
- fluid from an individual not having the disorder.

corresponding to this gene are useful for study, diagnosis of metabolic and immune disorders. The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 66

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HSYASIQSDDLWDSFNEVTNQTLDVKRMMKTWTLQKGFPLVTVQKKGKELFIQQERFFLNMKcomprise the amino acid sequence: EMFDSLSYFKGSSLLLMLKTYLSEDVFQHAVYLYLHN 1. (See Accession Nos. 2209276 and d1010078.) Preferred polypeptide fragments This translated product of this gene was recently identified as oxytocinase splice variant PEIQPSDTRYM (SEQ ID NO: 631). Also preferred are polynucleotide fragments encoding this polypeptide fragment.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 67

30 cells, and to lesser extent in primary dendritic cells and adipose tissue. This gene is expressed primarily in hemopoetic cells, particularly apoptotic T-

ઝ conditions: hemopoetic diseases including cancer and general immune disorders. reagents for differential identification of apoptotic T-cells, primary denritic cells, and providing immunological probes for differential identification of the tissue(s) or cell adipose tissue present in a biological sample and for diagnosis of diseases and Similarly, polypeptides and antibodies directed to these polypeptides are useful in Therefore, polynucleotides and polypeptides of the invention are useful as

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gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an and cell types (e.g., hematopoietic cells, and cancerous and wounded tissues) or bodily type(s). For a number of disorders of the above tissues or cells, particularly of the oral gene at significantly higher or lower levels may be routinely detected in certain tissues and intestinal mucosa as well as hemopoetic and immune systems, expression of this fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard individual not having the disorder.

corresponding to this gene are useful for treatment of diseases of the immune system, The tissue distribution indicates that polynucleotides and polypeptides including cancer, hemopoetic and infectious diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 68

This gene is expressed primarily in kidney cortex and to a lesser extent in infant

Therefore, polynucleotides and polypeptides of the invention are useful as brain, heart, uterus, and blood. 12

reagents for differential identification of kidney tissue present in a biological sample and fibrosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell for diagnosis of diseases and conditions: soft tissue cancer, inflammation, kidney

- relative to the standard gene expression level, i.e., the expression level in healthy tissue wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, lower levels may be routinely detected in certain tissues and cell types (e.g., kidney, type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and endocrines systems, expression of this gene at significantly higher or brain, and other nervous tissue, heart, uterus, and blood cells, and cancerous and or bodily fluid from an individual not having the disorder. ೧ 22
- corresponding to this gene are useful for study and treatment of cancer and fibroses. The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 69

The translation product of this gene shares strong sequence homology with vertebrate and invertebrate protein tyrosine phosphatases.

progenitors and to a lesser extent in infant brain, adipocytes, and several hematopoietic This gene is expressed primarily in endometrial tumors, melanocytes, myeloid stem cells. 35

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reagents for differential identification of transformed hematopoietic and epithelial cells Therefore, polynucleotides and polypeptides of the invention are useful as present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, of skin and endometrium, leukemia. Similarly,

- immunological probes for differential identification of the tissue(s) or cell type(s). For a hemopoietic systems, expression of this gene at significantly higher or lower levels may bone marrow, adipocytes, hematopoietic cells, and brain and other tissue of the nervous be routinely detected in certain tissues and cell types (e.g., endometrium, melanocytes, the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e., system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an polypeptides and antibodies directed to these polypeptides are useful in providing number of disorders of the above tissues or cells, particularly of the nervous and Š 2 15
- indicate that polynucleotides and polypeptides corresponding to this generate useful for The tissue distribution and sequence similarity with tyrosine phosphatases study and treatment of cancer and hematopoietic disorders.

FEATURES OF PROTEIN ENCODED BY GENE NO: 70 ន

This gene is expressed primarily in osteoclastoma, breast, and infant brain and to a lesser extent in various fetal and transformed bone, ovarian, and neuronal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

- polypeptides are useful in providing immunological probes for differential identification such a disorder, relative to the standard gene expression level, i.e., the expression level biological sample and for diagnosis of diseases and conditions: degenerative conditions cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and skeletal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell lypes (e.g., Juid or spinal fluid) or another tissue or cell sample taken from an individual having of the brain and skeleton. Similarly, polypeptides and antibodies directed to these bone, mammary tissue, and brain and other tissue of the nervous system; and 23 93
- in healthy tissue or bodily fluid from an individual not having the disorder. 35

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study and treatment of degenerative, neurological and skeletal disorders.

5 FEATURES OF PROTEIN: ENCODED BY GENE NO: 71

This gene was originally cloned from tumor cell lines. Recently another group has also cloned this gene, calling it the human malignant melanoma metastasis-suppressor (KiSS-1) gene. (See Accession No. U43527.) Preferred polypeptide fragments comprise the amino acid sequence: LEKVASVGNSRPTGQQLESLGLLA (SEQ ID NO: 632); VHREEASCYCQAEPSGDL (SEQ ID NO: 633); RPALRQAGGGTREPRQKRWAGL (SEQ ID NO: 634); and AVNFRPQRSQSM (SEQ ID NO: 635). Any frame shifts can easily be resolved using known molecular biology techniques.

This gene is expressed primarily in many types of carcinomas and to a lesser extent in many normal organs.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissues(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer particularly melanomas, and other hyperproliferative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in
- providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of transformed organ tissue, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to
- another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. As a tumor suppressor gene, increase amounts of the polypeptide can be used to treat patients having a particular cancer.

The tissue distribution indicates that this gene and the translated product is useful for diagnosing and study of cancer.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 72

This gene is expressed primarily in striatum and to a lesser extent in adipocytes and hemangioperiocytoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of striatal cells present in a biological sample and for diagnosis of diseases and conditions: neurological, fat and lysosomal storage

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diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., striatal tissue, adipocytes, and vascular tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis, study and treatment of neurodegenerative and growth disorders.

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 73

This gene is expressed primarily in bone marrow stromal cells and to a lesser extent in smooth muscle, testes, endothelium, and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of bone marrow present in a biological sample and for diagnosis of diseases and conditions: connective tissue and hematopoietic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal and hematopoietic systems, expression of this gene at significantly higher or

25 lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, stromal cells, smooth muscle, testes and other reproductive tissue, endothelium, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis, and treatment of connective tissue and blood diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 74

This gene is expressed primarily in brain, fetal liver and lung and to a lesser extent in retina, spinal chord, activated T-cells and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of brain and regenerating liver present in a biological sample and for diagnosis of diseases and conditions: CNS and spinal chord injuries, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

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higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, liver, pulmonary tissue, blood cells, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study and treatment of hematopoietic and neurological conditions.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 75

The translation product of this gene shares sequence homology with GTP binding proteins (intracellular).

This gene is expressed primarily in bone marrow, brain, and melanocytes and to a lesser extent in various endocrine and hematopoietic tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hematopietic and nervous system conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, melanocytes, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder,

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relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to nucleotide binding factors indicate that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis, and treatment of brain degenerative, skin and blood diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 76

This gene is expressed primarily in activated T-cells and to a lesser extent in retina, brain, and fetal bone.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of activated T-cells and developing brain present in a biological sample and for diagnosis of diseases and conditions: immune deficiencies and skeletal and neuronal growth disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous, immune, and skeletomuscular sustems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, brain and other tissue of the nervous system, retinal tissue, and bone, and cancerous and wounded tissues) or

bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis, study and treatment of cancer, urogenital, and brain degenerative diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 77

This gene is expressed primarily in fetal liver, activated monocytes, osteoblasts and to a lesser extent in synovial, brain, and lymphoid tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of myeloid and lymphoid present in a biological sample and for diagnosis of diseases and conditions: inflammation, immune deficiencies, cancer. Similarly, polypeptides and antibodies directed to these

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and skeleton, expression of this gene at significantly

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higher or lower levels may be routinely detected in certain tissues and cell types (e.g. expression level, i.e., the expression level in healthy tissue or bodily fluid from an taken from an individual having such a disorder, relative to the standard gene individual not having the disorder. serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., liver, blood cells, bone, synovial tissue, brain and other tissue of the nervous system

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and mesenchymal cancers and nervous system diseases. corresponding to this gene are useful for study, diagnosis, and treatment of lymphoid The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO:

polyprotein precursor which is thought to be important in DNA repair and replication The translation product of this gene shares sequence homology with polymerase

S and tumor cell lines This gene is expressed primarily in infant brain and to a lesser extent in tumors

not limited to, especially of the neural system and developing organs. Similarly, expression of this gene at significantly higher or lower levels may be routinely detected biological sample and for diagnosis of diseases and conditions which include, but are number of disorders of the above tissues or cells, particularly of the neural system immunological probes for differential identification of the tissue(s) or cell type(s). For a polypeptides and antibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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25 in certain (e.g., brain and other tissue of the nervous system, and cancerous and or bodily fluid from an individual not having the disorder wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder,

diagnosis and treatment of cancers especially of the neural system and developing indicate that polynucleotides and polypeptides corresponding to this gene are useful for The tissue distribution and homology to polymerase polyprotein precursor

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႘ FEATURES OF PROTEIN ENCODED BY GENE NO:

extent in brain. This gene is expressed primarily in muscle and endothelial cells and to a lesser

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providing immunological probes for differential identification of the tissue(s) or cell vascular system, expression of this gene at significantly higher or lower levels may be type(s). For a number of disorders of the above tissues or cells, particularly of the Similarly, polypeptides and antibodies directed to these polypeptides are useful in biological sample and for diagnosis of diseases and conditions: vascular diseases reagents for differential identification of the tissue(s) or cell type(s) present in a nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, routinely detected in certain (e.g., muscle, endothelial cells, brain and other tissue of the Therefore, polynucleotides and polypeptides of the invention are useful as

5 corresponding to this gene are useful for treatment and diagnosis of disorders of the plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e. the expression level in healthy tissue or bodily fluid from an individual not having the The tissue distribution indicates that polynucleotides and polypeptides

5 vascular and neural system including cardiovascular and endothelial

FEATURES OF PROTEIN ENCODED BY GENE NO: 80

This gene is expressed primarily in placenta and to a lesser extent in fetal liver

23 20 immunological probes for differential identification of the tissue(s) or cell type(s). For a reagents for differential identification of the tissue(s) or cell type(s) present in a of this gene at significantly higher or lower levels may be routinely detected in certain number of disorders of the above tissues or cells, particularly of developmental and disorder of the haemopoietic system, fetal liver and placenta. Similarly biological sample and for diagnosis of diseases and conditions: developmental disorders polypeptides and antibodies directed to these polypeptides are useful in providing tissues and cell types (e.g., placenta and liver, and cancerous and wounded tissues) or disorders and disorder of the haemopoietic system, fetal liver and placenta, expression Therefore, polynucleotides and polypeptides of the invention are useful as

ઝ bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the fluid from an individual not having the disorder.

corresponding to this gene are useful for diagnosis and treatment of developmental disorders and disorders of the haemopoietic system, fetal liver and placenta The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 81

This gene is expressed primarily in bone marrow, placenta and tissues and organs of the hematopoietic system.

polypeptides are useful in providing immunological probes for differential identification relative to the standard gene expression level, i.e., the expression level in healthy tissue of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, bone and hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, biological sample and for diagnosis of diseases and conditions: disorders of the bone Therefore, polynucleotides and polypeptides of the invention are useful as and haemopoietic system. Similarly, polypeptides and antibodies directed to these types (e.g., bone marrow, placenta, and hematopoietic cells, and cancerous and reagents for differential identification of the tissue(s) or cell type(s) present in a or bodily fluid from an individual not having the disorder. 2

corresponding to this gene are useful for diagnosis and treatment of disorders of the The tissue distribution indicates that polynucleotides and polypeptides

immune, bone and hematopoietic system ន

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FEATURES OF PROTEIN ENCODED BY GENE NO: 82

The translation product of this gene shares sequence homology with secretory carrier membrane protein which is thought to be important in protein transport and export. Any frame shifts in coding sequence can be easily resolved using standard molecular biology techniques. Another group recently cloned this gene, calling it SCAMP. (See Accession No. 2232243.) 25

This gene is expressed primarily in prostate, breast and spleen, and to a lesser extent in several other tissues and organs.

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, significantly higher or lower levels may be routinely detected in certain tissues and cell biological sample and for diagnosis of diseases and conditions: disorders of the breast Therefore, polynucleotides and polypeptides of the invention are useful as particularly disorders of the breast prostate and spleen, expression of this gene at reagents for differential identification of the tissue(s) or cell type(s) present in a prostate and spleen. Similarly, polypeptides and antibodies directed to these 33 30

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or types (e.g., prostate, mammary tissue, and spleen, and cancerous and wounded fluid from an individual not having the disorder.

indicate that polynucleotides and polypeptides corresponding to this gene are useful for The ussue distribution and homology to secretory carrier membrane protein diagnosis and treatment of disorders of the breast, prostate and spleen.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 83 2

This gene is expressed primarily in developing organs and tissue like placenta and infant brain and to a lesser extent in developed organs and tissue like cerebellum

routinely detected in certain tissues and cell types (e.g., placenta, heart, brain and other biological sample and for diagnosis of diseases and conditions: neurological diseases. sample taken from an individual having such a disorder, relative to the standard gene neural system, expression of this gene at significantly higher or lower levels may be providing immunological probes for differential identification of the tissue(s) or cell issue of the nervous system, and cancerous and wounded tissues) or bodily fluids expression level, i.e., the expression level in healthy tissue or bodily fluid from an (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Similarly, polypeptides and antibodies directed to these polypeptides are useful in Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. 15 ន 25

corresponding to this gene are useful for treatment and diagnosis of diseases of the The tissue distribution indicates that polynucleotides and polypeptides neural system including neurological disorders and cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 84

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The translation product of this gene shares sequence homology with ATPase 6 in Trypanosoma brucei which is thought to be important in metabolism

This gene is expressed primarily in tumor and fetal tissues and to a lesser extent in melanocytes, kidney cortex, monocytes and ovary.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions: metabolism disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., fetal tissues, melanocytes, kidney, blood cells, ovary and other tissue of the reproductive system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

The tissue distribution and homology to ATPase indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of metabolism disorders, especially in fetal and tumor tissue growth.

FEATURES OF PROTEIN ENCODED BY GENE NO: 85

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fluid from an individual not having the disorder

The translation product of this gene shares sequence homology with the immunoglobulin superfamily of proteins which are known to be important in immune response and immunity.

20 This gene is expressed primarily in stromal cells, colon cancer, lung, amygdala, melanocyte and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of stromal cell development and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the stromal cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., stromal cells,

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colon, lung, amygdala, and melanocytes, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to immunoglobulin indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of immune system disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 86

The translation product of this gene shares sequence homology with transcription iniation factor eIF-4 gamma which is thought to be important in gene transcription

This gene is expressed primarily in tumor tissues

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumorigenesis.

- providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly in tumor tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in
- The tissue distribution and homology to transcription iniation factor eIF-4
 20 gamma indicate that polynucleotides and polypeptides corresponding to this gene are
 useful for gene regulation in tumorigenesis.

healthy tissue or bodily fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 87

The translation product of this gene shares sequence homology at low level in prolines with secreted basic proline-rich peptide II-2 which is thought to be important in protein structure or inhibiting hydroxyapatite formation in vitro.

This gene is expressed primarily in endometrial tumor and fetal lung

Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions: endometrial tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the muscular/skeletal and reproductive systems, expression of this gene at significantly

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35 higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample

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taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to secreted basic proline-rich peptide II-2 indicate that polynucleotides and polypeptides corresponding to this gene are useful for inhibiting hydroxyapatite formation or establishing cell/tissue structure.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 88

This gene is expressed primarily in: amniotic cells inducted with TNF in culture; and to a lesser extent in colon tissue from a patient with Crohn's Disease; parathyroid turnor; activated T-cells; cells of the human Caco-2 cell line; adenocarcinoma; colon; corpus colosum; fetal kidney; pancreas tumor; fetal brain; early stage brain, and anergic

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system;

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c.g., tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain (e.g., amniotic cells, colon, kidney, pancreas, parathyroid, brain and other tissue of the nervous system, blood cells, hematopoietic cells, liver, spleen, bone, testes and other reproductive tissue, brain and other tissue of the nervous system, and epithelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product of this clone is useful for modulating tumorigenesis and other immune system conditions such as disorders in immune response.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 89

This gene is expressed primarily in fetal liver/spleen and hematopoietic cells and

to a lesser extent in brain, osteosarcoma, and testis tumor.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions: leukemia and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

significantly of the hematopoictic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, liver, spleen, bone, testes, and other reproductive tissue, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or 10 another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopoietic and immune disorders.

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fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 90

The translation product of this gene shares weak sequence homology with mouse Gcap1 protein which is developmentally regulated in brain.

This gene is expressed primarily in infant and adult brain and fetal liver/spleen and to a lesser extent in smooth muscle, T cells, and a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological or hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous, hematopoietic, immune, and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain

itssues and cell types (e.g., brain and other tissue of the nervous system, blood cells, liver, spleen, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and its homology to Gcap1 protein indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatubg and diagnosis of disorders in neuronal, hematopoietic, immune, and endocrine systems.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 91

This gene is expressed primarily in brain and hematopoietic cells and to a lesser extent in tumor tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a revous.

- hematopoietic, immune systems and tumorigenesis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the in nervous, hematopoietic, immune
- detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of disorders in the nervous, hematopoietic, and immune systems.

FEATURES OF PROTEIN ENCODED BY GENE NO: 92

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The translation product of this gene shares sequence homology with neuroendocrine-specific protein A which is thought to be important in neurologic

30 This gene is expressed primarily in brain tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions; neural disorders and degeneration disease. Similarly, polypeptides and antibodies directed to these

35 polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central or peripheral nervous systems, expression of this gene at

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significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level

in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to neuroendocrine-specific protein A

In the tissue distribution and notificing to incurrence of this gene are useful for indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment or diagnosis of neural disorders and degeneration disease.

FEATURES OF PROTEIN ENCODED BY GENE NO: 93

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The translation product of this gene shares sequence homology with collagenlike protein and prolin-rich protein which are thought to be important in connective tissue function and tissue structure.

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This gene is expressed primarily in fetal liver/spleen and brain tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neuronal or hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and hematopoietic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial

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The tissue distribution and homology to collagen-like protein and proline-rich groteins indicate that polynucleotides and polypeptides corresponding to this gene are useful for supporting brain and hematopoietic tissue function and diagnosis and treatment of disorders in these functions.

in healthy tissue or bodily fluid from an individual not having the disorder.

fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level

FEATURES OF PROTEIN ENCODED BY GENE NO: 94

This gene is expressed primarily in embryonic tissues and tumor tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions which include, but are not limited to.. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system (e.g., tumors), expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancer.

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15 FEATURES OF PROTEIN ENCODED BY GENE NO: 95

This gene is expressed primarily in brain tumor, placenta, and melanoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: brain tumor or melanoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain or melanocytes, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, placenta, and melanocytes, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

30 The tissue distribution indicates that the translation product of this gene is useful in the diagnosis and treatment of brain tumors and melanoma.

FEATURES OF PROTEIN ENCODED BY GENE NO: 96

The translation product of this gene shares sequence homology with a yeast membrane protein, SUR4, which encodes for APA1 that acts on a glucose-signaling pathway that controls the expression of several genes that are transcriptionally regulated by glucose.

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This gene is expressed primarily in fetal liver, and to a lesser extent in placenta the set tissue

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of fetal liver or defects of glucose-regulated ATPase activities in tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal immune/hematopoietic system,

in certain tissues and cell types (e.g., liver, placenta, and mammary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual or having the disorder.

The tissue distribution and homology to yeast SUR4 membrane prolein indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of defects of fetal liver or defects of glucose-regulated ATPase

FEATURES OF PROTEIN ENCODED BY GENE NO: 97

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This gene is expressed primarily in fetal liver, brain, and amniotic fluid.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of the fetal immune system and adult brain. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal immune system and adult brain, expression of this gene at

significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., anniotic fluid, serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder

The tissue distribution indicates that the protein product of this clone is useful for detecting defects of the fetal immune and hematopoietic systems since fetal liver is

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the predominant organ responsible for hematopoiesis in the fetus. In addition, the gene product of this gene is thought to be useful for detecting certain neurological defects of

FEATURES OF PROTEIN ENCODED BY GENE NO: 98

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The translation product of this gene shares sequence homology with an yolk protein precursor, Vitellogenin which is thought to be important in binding lipids such as phosvitin.

This gene is expressed primarily in amniotic cells and fetal liver.

5 20 15 biological sample and for diagnosis of diseases and conditions: defects in amniotic reagents for differential identification of the tissue(s) or cell type(s) present in a for differential identification of the tissue(s) or cell type(s). For a number of disorders cells, fetal liver development and the fetal immune system. Similarly, polypeptides and urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, levels may be routinely detected in certain tissues and cell types (e.g., amniotic cells, state is likely, e.g., immunel, expression of this gene at significantly higher or lower of the above tissues or cells, particularly of the [insert system where a related disease antibodies directed to these polypeptides are useful in providing immunological probes the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e., Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution and homology to vitellogenin indicate that the protein product of this clone is useful for treatment and diagnosis of defects in amniotic cells, fetal liver development and the fetal immune system.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 99

This gene is expressed primarily in placenta, endometrial tumor, osteosarcoma and stromal cells.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumor of the endometrium or bone, and osteosarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the obstetric system (e.g. placenta,

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endometrium) and the bones, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, endometrium, bone, and stromal cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tumors and 10 abnormalities of the endometrium, and the bones because of its abundance in the aforementioned tissues...

FEATURES OF PROTEIN ENCODED BY GENE NO: 100

This gene is expressed primarily in hepatocellular tumor.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hepatocellular tumor. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the liver expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of hepatocellular cancer because of its abundant expression in this tissue.

FEATURES OF PROTEIN ENCODED BY GENE NO: 101

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This gene is expressed primarily in Corpus Colosum, fetal lung and infant

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of the Corpus Colosum or defects of the fetal lung. Similarly, polypeptides and antibodies directed to

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brain

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these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Corpus Colosum and brain in general, and fetal lung, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., lung, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of defects of the Corpus Colosum and brain in general, and defects of fetal lung.

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15 FEATURES OF PROTEIN ENCODED BY GENE NO: 102

This gene is expressed primarily in T cells and stromal cells, and to a lesser extent in adrenal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of T cell immunity and stromal cell development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell

ussues or cells, particularly of the infiniture system, expression of this gold at tissues and cell significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, stromal cells, and adrenal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of defects of T cell immunity and stromal cell development because of its abundant expression in these tissues.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 103

This gene is expressed primarily in infant brain and placenta.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of the brain and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, especially brain, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and placenta, cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that the protein product of this clone is useful for detecting defects of the brain, especially in young children.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 105

This gene is expressed primarily in human osteoclastoma and to a lesser extent in human pancreas tumor.

another tissue or cell sample taken from an individual having such a disorder, relative to immunological probes for differential identification of the tissue(s) or cell type(s). For a expression of this gene at significantly higher or lower levels may be routinely detected the standard gene expression level, i.e., the expression level in healthy tissue or bodily in certain tissues and cell types (e.g., bone and pancreas, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or biological sample and for diagnosis of diseases and conditions which include, but are number of disorders of the above tissues or cells, particularly in transformed tissues, Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and antibodies directed to these polypeptides are useful in providing not limited to, cancer particularly osteoclastoma and pancreatic tumor. Similarly, reagents for differential identification of the tissue(s) or cell type(s) present in a fluid from an individual not having the disorder. 22 ಜ 2

The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of some types of tumors, particularly pancreatic cancer and osteoclastoma.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 106

This gene is expressed primarily in fetal liver/spleen, and to a lesser extent in vated T-Cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the

- immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the
- 15 expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of immune disorders.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 107

This gene is expressed primarily in human embryo and to a lesser extent in spleen and chronic lymphocytic leukemia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: leukemia. Similarly,

biological sample and for diagnosis of diseases and conditions: leukemia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune or hemopoietic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, spleen, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution indicates that the protein product of this clone is useful for the diagnosis and treatment of leukemia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 108

This gene is expressed primarily in placenta, and to a lesser extent in early stage human brain and in lung.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: fetal developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly in fetal and amniotic tissue, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, brain and other tissue of the nervous system, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that the protein product of this is useful for production of growth factor(s) associated with fetal development. Preferred polypeptides comprise the full-length polypeptide shown in the sequence listing, truncated however, at the amino terminus and beginning with QTIE.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 109

This gene is expressed primarily in fetal spleen, and to a lesser extent in B-Cell

25 lymphoma and T-Cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., spleen and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that the protein product of this clone is useful for the treatment and diagnosis of human lymphomas.

FEATURES OF PROTEIN ENCODED BY GENE NO: 110

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The translation product of this gene shares sequence homology with sarcoma amplified sequence (SAS), a tetraspan receptor which is thought to be important in malignant fibrous histiocytoma and liposarcoma.

This gene is expressed primarily in human osteoclastoma, and to a lesser extent in pineal gland and infant brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: malignant fibrous histiocytoma and liposarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone, pineal gland, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to sarcoma amplified sequence (SAS) indicate that the protein product of this clone is useful for treatment of, osteosarcoma, malignant fibrous histiocytoma and liposarcoma and related cancers, particularly sarcomas.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 111

The translation product of this gene shares sequence homology with 6.8K

proteolipid protein, mitochondrial - bovine.

This gene is expressed primarily in Wilm's tumor and to a lesser extent in

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cerebellum and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Wilm's tumor. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunblogical probes for differential identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the immune or renal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

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The tissue distribution and homology to 6.8K proteolipid protein indicate that 10 the protein product of this clone is useful for diagnostic and therapeutics associated with tumors, particularly Wilm's tumor disease.

individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 112

This gene is expressed primarily in embryonic tissue and to a lesser extent in osteoblasts, endothelial cells, macrophages (GM-CSF treated), and bone marrow.

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osteoblasts, endothelial cells, macrophages (GM-CSF treated), and bone marrow. Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in

- type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, bone, endothelial cells, blood cells and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., enum plasma prime synowial fluid or spinal fluid) or another
 - bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides

corresponding to this gene are useful for treatment or diagnosis of immune disorders.

Preferred polypeptides encoded by this gene comprise the following amino acid
sequence: MITDVQLAIFANMLGVSLFLLVVLYHYVAVNNPKKQE (SEQ ID NO: 636).

FEATURES OF PROTEIN ENCODED BY GENE NO: 113

35 This gene is expressed primarily in hepatocellular tumor, and to a lesser extent in fetal liver/spleen.

urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma particularly of the hepatic system, expression of this gene at significantly higher or biological sample and for diagnosis of diseases and conditions: tumors, particularly individual having such a disorder, relative to the standard gene expression level, i.e., lower levels may be routinely detected in certain tissues and cell types (e.g., liver, and hepatocellular tumors. Similarly, polypeptides and antibodies directed to these reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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for diagnosis and treatment of tumors, particularly hepatocellular tumors. The tissue distribution indicates that the protein product of this clone is useful

the expression level in healthy tissue or bodily fluid from an individual not having the

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FEATURES OF PROTEIN ENCODED BY GENE NO: 114

colleagues (Nature 389, 300-306 (1997)). In addition, the predicted translation product apoptosis. The sequence of this gene has since been published by Polyak and EI24 which is also thought to be important in p53 mediated apoptosis. of this contig exhibits very high sequence homology with a murine gene denoted as identity with the human Pig8 gene which is thought to be important in p53 mediated The translation product of this gene exhibits a very high degree of sequence

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lesser extent in bone marrow, fetal liver, and prostate. This gene is expressed primarily in infant brain and activated T-cells and to a

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reagents for differential identification of the tissue(s) or cell type(s) present in a not limited to, and tissue damage by radiation and anti-cancer drugs. Similarly, biological sample and for diagnosis of diseases and conditions which include, but are Therefore, polynucleotides and polypeptides of the invention are useful as

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ઝ polypeptides and antibodies directed to these polypeptides are useful in providing nervous system, blood cells, bone marrow, liver, and prostate, and cancerous and routinely detected in certain tissues and cell types (e.g., brain and other tissue of the number of disorders of the above tissues or cells, particularly of the nervous and immunological probes for differential identification of the tissue(s) or cell type(s). For a wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal immune systems, expression of this gene at significantly higher or lower levels may be fluid) or another tissue or cell sample taken from an individual having such a disorder,

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or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue

also be useful in the treatment of hematopoietic disorders and in boosting numbers of is upregulated in cells undergoing such treatment where p53 was overexpressed. It may etoposide, hydroperoxycyclophosphamide, and X-irradiation, since this protein product preventing apoptosis in patients being treated with anti-oncogenic drugs such as indicate that polynucleotides and polypeptides corresponding to this gene are useful for hematopoietic stem cells by interfering with the apoptosis of progenitor cells. The The tissue distribution and homology to human Pig8 and murine E124 genes

5 5 mature polypeptide is predicted to comprise the following amino acid sequence: SSYIISGCLFSILFPLFIISANEAKTPGKAYLFQLRLFSLVVFLSNRLFHKTVYLQSALSSSTSAEK EPRIVSRIFQCCAWNGGVFWFSLLLFYRVFIPVLQSVTARIIGDPSLHGDVWSWLEFFLTSIFSA FPSPHPSPAKLKATAGH (SEQ ID NO: 637). Accordingly, polypeptides comprising the FPIHLVGQLVSLLHMSLLYSLYCFEYRWFNKGIEMHQRLSNIERNWPYYFGFGLPLAFLTAMQ LWYLPLFYLSKYVNAIWFQDIADLAFEYSGRKPHPFPSYSKIIADMLFNLLLQALFLIQGMFYSI eemadsvktflqdlargikdsiwgictiskldariqqkreeqrrrrassvlaqrraqsierkqes foregoing amino acid sequence are provided as are polynucleotides encoded such

FEATURES OF PROTEIN ENCODED BY GENE NO: 115

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multiple sclerosis. This gene is expressed primarily in stromal cells and to a lesser extent in

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

- 25 30 biological sample and for diagnosis of diseases and conditions: affecting the nervous routinely detected in certain tissues and cell types (e.g., stromal cells and cancerous and in providing immunological probes for differential identification of the tissue(s) or cell system. Similarly, polypeptides and antibodies directed to these polypeptides are useful type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal
- 35 and other autoimmune diseases. corresponding to this gene are useful for treatment and diagnosis of multiple sclerosis The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 116

This gene is expressed primarily in the gall bladder

polypeptides are useful in providing immunological probes for differential identification biological sample and for diagnosis of diseases and conditions: gall stones or infection of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, Therefore, polynucleotides and polypeptides of the invention are useful as of the digestive system. Similarly, polypeptides and antibodies directed to these reagents for differential identification of the tissue(s) or cell type(s) present in a S

types (e.g., gall bladder and tissue of the digestive system, and cancerous and wounded another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily significantly higher or lower levels may be routinely detected in certain tissues and cell tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or particularly of the digestive system or renal system, expression of this gene at fluid from an individual not having the disorder. 2 2

corresponding to this gene are useful for possible prevention of digestive disorders where there may be a lack of digestive enzymes produced or in the detection and The tissue distribution indicates that polynucleotides and polypeptides

possible prevention of gall stones. 8

FEATURES OF PROTEIN ENCODED BY GENE NO: 117

The translation product of this gene shares sequence homology with dystrophin This gene is expressed primarily in placenta and to a lesser extent in fetal brain gene which is thought to be important in building and maintenance of muscles.

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Therefore, polynucleotides and polypeptides of the invention are useful as and fetal liver, and spleen.

the above lissues or cells, particularly of the skeletal muscle system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues differential identification of the tissue(s) or cell type(s). For a number of disorders of liver, and spicen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, Duchenne and Becker's muscular dystropies. Similarly, polypeptides and antibodies and cell types (e.g., placenta, brain and other tissue of the nervous system, muscle, biological sample and for diagnosis of diseases and conditions: muscular dystropy, reagents for differential identification of the tissue(s) or cell type(s) present in a directed to these polypeptides are useful in providing immunological probes for 35

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plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from

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an individual having such a disorder telative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

related the degenerative myopathies that are characterized by the weakness and atrophy polynucleotides and polypeptides corresponding to this gene are useful for diseases of muscles without neural degradation; such as Duchenne and Becker's muscular The tissue distribution and homology to the dystrophin gene indicate that S

FEATURES OF PROTEIN ENCODED BY GENE NO: 118 2

dystropies.

This gene is expressed primarily in olfactory tissue and to a lesser extent in

Therefore, polynucleotides and polypeptides of the invention are useful as

- polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the connective tissue, expression of this gene at significantly higher or diseases; chondrosarcoma. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: connective tissue reagents for differential identification of the tissue(s) or cell type(s) present in a 12 ន
- an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from tissue and cartilage, and cancerous and wounded tissues) or bodily fluids (e.g., serum, the expression level in healthy tissue or bodily fluid from an individual not having the lower levels may be routinely detected in certain tissues and cell types (e.g., olfactory

corresponding to this gene are useful for tumors of connective tissues, osteparthritis The tissue distribution indicates that polynucleotides and polypeptides and the treatment and diagnosis of chondrosarcoma.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 119 9

This gene is expressed primarily in Activated Neutrophils and to a lesser extent in fetal spleen, and CD34 positive cells from cord blood.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

hematopoiesis and inflammation. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: allergies, defects in these polypeptides are useful in providing immunological probes for differential 35

identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and hematopoiesis system the, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for reducing the allergic effects felt by allergy suffers by neutralizing the activity of the immune system, especially since neutrophils are abundant in persons suffering from allergies and other inflammatory conditions.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 120

The translation product of this gene shares sequence homology with poly A binding protein Π which is thought to be important in RNA binding for transcription of RNA to DNA

This gene is expressed primarily in colon and to a lesser extent in brain and

20 25 မ number of disorders of the above tissues or cells, particularly of the immune and polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: colon cancer. Similarly, reagents for differential identification of the tissue(s) or cell type(s) present in a routinely detected in certain tissues and cell types (e.g., colon, tissue and cells of the digestive system, expression of this gene at significantly higher or lower levels may be or bodily fluid from an individual not having the disorder relative to the standard gene expression level, i.e., the expression level in healthy tissue immune system, and brain or other tissue of the nervous system, and cancerous and immunological probes for differential identification of the tissue(s) or cell type(s). For a fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal Therefore, polynucleotides and polypeptides of the invention are useful as The tissue distribution and homology to poly A binding protein II indicate that

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FEATURES OF PROTEIN ENCODED BY GENE NO: 121

The translation product of this gene shares sequence homology with thymidine diphosphoglucose 4.6 dehydrase which is thought to be important in the metabolism of sugar.

This gene is expressed primarily in fetal liver and spleen and to a lesser extent in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diabetes. Similarly,

- immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
- The tissue distribution and homology to thyrnidine diphospoglucose 4.6 dehydrase indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of persons with diabetes since it appears that this protein is needed in the metabolism of sugar in to its more basic components.

25 FEATURES OF PROTEIN ENCODED BY GENE NO: 122

The translation product of this gene shares sequence homology with ceruloplasmin which is thought to be important in the metabolism and transport of iron and copper. Ceruloplasmin also contains domains with homology to clotting factors V and VIII. Defects in the circulating levels of ceruloplasmin (aceruloplasminemia) have

been associated with certain disease conditions such as Wilson disease, and the accompanying hepatolenticular degeneration.

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This gene is expressed primarily in brain and retina and to a lesser extent in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases marked by defects in iron metabolism; accruloplasminemia not characterized by defects in the

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and treatment of colon cancer and other disorders of the digestive system.

polynucleotides and polypeptides corresponding to this gene are useful for detection

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known ceruloplasmin gene locus; nonclassical Wilson disease; movement disorders; and tumors derived from a brain tissue origin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, retina, and nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, retinal tissue, and endothelial cells, and cancerous and wounded tissues) or bodily

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fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ceruloplasmin indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of patients with aceruloplasminemia, or other defects in iron and/or copper metabolism. Mutations in this locus could also be diagnostic for patients currently experiencing or predicted to experience aceruloplasminemia.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 123

This gene is expressed primarily in brain and B cell lymphoma and to a lesser extent in fetal liver and spleen.

may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of immunological probes for differential identification of the tissue(s) or cell type(s). For a the nervous system, blood cells, liver, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another hematopoietic system, expression of this gene at significantly higher or lower levels lissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: B cell lymphoma; tumors and diseases of the brain and/or spleen; hematopoietic defects. Similarly, reagents for differential identification of the tissue(s) or cell type(s) present in a number of disorders of the above tissues or cells, particularly of the brain and fluid from an individual not having the disorder. 33 23 2

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of disorders in neuronal,

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hematopoietic, and immune systems. It could potentially be useful for neurodegenerative disorders and neuronal and/or hematopoietic cell survival or proliferation.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 124

This gene is expressed primarily in osteoclastoma, dermatofibrosarcoma, and B cell lymphoma and to a lesser extent in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer in particular osteoclastoma, dermatofibrosarcoma, and B cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of

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15 the bone, immune, and circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone, epidermis, blood cells, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancers and lymphoma; osteoporosis; and the control of cell proliferation and/or differentiation.

fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 125

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This gene is expressed primarily in immune tissues and hematopoielic cells, particularly in activated T cells and neutrophils, spleen, and fetal liver, and to a lesser extent in infant adrenal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects in T cell activation; hematopoietic disorders; tumors of a hematopoietic and/or adrenal gland origin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and/or endocrine systems, expression of this gene at significantly higher

or lower levels may be routinely detected in certain tissues and cell types (e.g., cells and tissues of the immune system, hematopoietic cells, blood cells, liver, and adrenal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune and/or hematopoietic disorders; diseases related to proliferation and/or differentiation of hematopoietic cells; defects in T cell and neutrophil activation and responsiveness; and endocrine and/or metabolic disorders, particularly of early childhood.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 126

This gene is expressed primarily in placenta and endothelial cells and to a lesser extent in melanocytes and embryonic tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors of an endothelial cell origin; angiogenesis associated with tumor development and metastasis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system and developing embryo, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, endothelial cells, melanocytes, and embryonic tissues, and cancerous and wounded tissues) or

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bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of developmental disorders; inhibition of angiogenesis; and vascular patterning.

35 FEATURES OF PROTEIN ENCODED BY GENE NO: 127

This gene is expressed primarily in endothelial cells and hematopoietic tissues, including spleen, tonsils, leukocytes, and both B- and T-cell lymphomas.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors of an endothelial cell and/or hematopoietic origin; leukemias and lymphomas. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and vascular systems expression of this gene at significantly higher or lower levels may be routinely detected

in certain tissues and cell types (e.g., endothelial cells, hematopoietic cells, spleen, 10 tonsils, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the manipulation of angiogenesis; the differentiation and morphogenesis of endothelial cells; the proliferation and/or differentiation of hematopoietic cells; and the commitment of hematopoietic cells to distinct cell lineages.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 128

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This gene is expressed primarily in kidney medulla and to a lesser extent in spleen from chronic myelogenous leukemia patients, prostate cancer, and some other tissues.

25 30 ઝ reagents for differential identification of the tissue(s) or cell type(s) present in a for differential identification of the tissue(s) or cell type(s). For a number of disorders biological sample and for diagnosis of diseases and conditions: tumors of a kidney of the above tissues or cells, particularly of the kidney and spleen, expression of this origin; chromic myelogenous leukemia; prostate cancer. Similarly, polypeptides and standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another and cell types (e.g., kidney, spleen, and prostate, and cancerous and wounded tissues gene at significantly higher or lower levels may be routinely detected in certain tissues antibodies directed to these polypeptides are useful in providing immunological probes fluid from an individual not having the disorder. Therefore, polynucleotides and polypeptides of the invention are useful as

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis and treatment of kidney disorders and cancer, particularly chromic myelogenous leukemia and prostate cancer. It may also be useful for the enhancement of kidney tubule regeneration in the treatment of acute renal failure.

FEATURES OF PROTEIN ENCODED BY GENE NO: 129

This gene is expressed primarily in adult and infant brain and to a lesser extent in mesenchymal or fibroblast cells, as well as tissues with a mesenchymal origin.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors of a brain and/or mesenchymal origin; neurodegenerative disorders; cancer; fibrosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing irmmunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and of mesenchymal cells and tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system and cancerous and wounded tissues) or bodily fluids

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20 (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the diagnosis of tumors of a brain and/or mesenchymal origin; neurodegenerative disorders; cancer; and fibrosis, based upon the expression of this gene within those tissues. Fibrosis is considered as mesenchymal cells and fibroblasts are the primary cellular targets involved in this pathological condition.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 130

This gene is expressed primarily in hepatocellular cancer and to a lesser extent in fetal tissues as well as testes tumor.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: liver cancer. Similarly, polypeptides and altibodies directed to these polypeptides are useful in providing

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immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, fetal tissue, and testes and other

- reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
- 10 The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of liver cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 131

This gene is expressed only in infant early brain.

- reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: development and diseases of the nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the
- The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating diseases of the brain in children and in treating nervous system disorders such as Alzheimer's disease, schizophrenia, dementia, depression, etc.

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standard gene expression level, i.e., the expression level in healthy tissue or bodily

fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 132

This gene is expressed primarily in brain and to a lesser extent in glioblastoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Alzheimer's disease,

schizophrenia, depression, mania, and dementia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating brain disorders such as Alzheimer's disease, schizophrenia, depression, mania, and dementia.

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healthy tissue or bodily fluid from an individual not having the disorder

15 FEATURES OF PROTEIN ENCODED BY GENE NO: 133

The translation product of this gene shares sequence homology with ribitol dehydrogenase of bacteria which is thought to be important in metabolism of sugars.

This gene is expressed primarily in macrophage and to a lesser extent in T-cell

lymphoma and lung.

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tissue destruction in inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ribitol dehydrogenase indicate that polynucleotides and polypeptides corresponding to this gene are useful for altering macrophage metabolism in diseases such as inflammation where macrophages are causing excess tissue destruction.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 13

This gene is expressed primarily in pancreatic tumor and to a lesser extent in synovial sarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to,. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine and connective tissue systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., pancreas, and synovial tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating and diagnosing various cancers.

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 135

This gene is expressed primarily in T cell lines such as Raji and to a lesser extent in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune system disorders and inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or

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30 lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating and diagnosing inflammatory diseases

such as rheumatoid arthritis, sepsis, inflammatory bowel disease, and psoriasis, as well as neutropenia.

FEATURES OF PROTEIN ENCODED BY GENE NO: 136

The translation product of this gene shares high sequence homology with SAR1 subfamily of GTP-binding proteins which is thought to be important in vesicular transport in mammalian cells.

This gene is expressed primarily in serum-stimulated smooth muscle cells and to a lesser extent in a T-cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases affecting vesicular transport. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the muscular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

polynucleotides and polypeptides corresponding to this gene are useful for gene therapy in treating the large number of diseases involved in defective vesicular transport within cells...

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individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 137

The translation product of this gene shares sequence homology with a protein found in C. elegans cosmid F25B5.

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This gene is expressed primarily in a fetal tissues and to a lesser extent in melanocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: abnormal fetal development, especially of the pulmonary system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

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for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal pulmonary system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., fetal tissue, pulmonary tissue, and melanocytes, and

5 cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of diseases affecting the pulmonary system, such as emphysema.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 138

This gene is expressed primarily in gall bladder and to a lesser extent in smooth

muscle.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: digestive system disease and gall bladder problems. Similarly, polypeptides and antibodies directed to these

- polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., gall bladder and tissue of the digestive system, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal
 - oranger and tissue of the digestive system, and smooth muscie, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating diseases of the digestive system.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 139

This gene is expressed primarily in placenta and to a lesser extent in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as
reagents for differential identification of the tissue(s) or cell type(s) present in a
biological sample and for diagnosis of diseases and conditions: abnormal fetal
development. Similarly, polypeptides and antibodies directed to these polypeptides are

tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids expression level, i.e., the expression level in healthy tissue or bodily fluid from an sample taken from an individual having such a disorder, relative to the standard gene (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell individual not having the disorder.

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5 corresponding to this gene are useful for treating and diagnosing abnormal fetal The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 140

2 ovary, prostate cancer, and activated monocytes. This gene is expressed primarily in smooth muscle and to a lesser extent in

biological sample and for diagnosis of diseases and conditions: hypertension and reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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atherosclerosis. Similarly, polypeptides and antibodies directed to these polypeptides muscle, ovary and other reproductive tissue, prostate, and blood cells, and cancerous are useful in providing immunological probes for differential identification of the lower levels may be routinely detected in certain tissues and cell types (e.g., smooth particularly of the circulatory system, expression of this gene at significantly higher or tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

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မ such as hypertension, atherosclerosis, etc. corresponding to this gene are useful for treating diseases of the circulatory system The tissue distribution indicates that polynucleotides and polypeptides

healthy tissue or bodily fluid from an individual not having the disorder.

disorder, relative to the standard gene expression level, i.e., the expression level in spinal fluid) or another tissue or cell sample taken from an individual having such a and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or

FEATURES OF PROTEIN ENCODED BY GENE NO: 141

33 and bone marrow This gene is expressed primarily in fetal spleen and to a lesser extent in placenta

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of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification types (e.g., spieen, placenta, bone marrow, and blood cells, and cancerous and significantly higher or lower levels may be routinely detected in certain tissues and cell particularly of the circulatory and pulmonary systems, expression of this gene at diseases affecting blood cells. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: anemia and other reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

5 or bodily fluid from an individual not having the disorder relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal

2 corresponding to this gene are useful for the generation of red and white blood cells and for the diagnosis of disease of these cells. The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 142

20 murine tetracycline/sugar transporter molecule recently reported by Matsuo and colleagues (Biochem. Biophys. Res. Commun. 238 (1), 126-129 (1997)). The predicted translation product of this contig is a human homolog of the

This gene is expressed primarily in synovium and to a lesser extent in

reagents for differential identification of the tissue(s) or cell type(s) present in a or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: rheumatoid arthritis and Therefore, polynucleotides and polypeptides of the invention are useful as

- ಅ the immune and lymphatic systems, expression of this gene at significantly higher or serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample tissue, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., lower levels may be routinely detected in certain tissues and cell types (e.g., synovial taken from an individual having such a disorder, relative to the standard gene
- 35 expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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diseases, such as rheumatoid arthritis, leukemia, neutropenia, inflammatory bowel corresponding to this gene are useful for treatment and diagnosis of inflammatory The tissue distribution indicates that polynucleotides and polypeptides disease, psoriasis, sepsis, and the like.

FEATURES OF PROTEIN ENCODED BY GENE NO: 143

This gene is expressed primarily in placenta and to a lesser extent in melanocyte, fetal liver and spleen, and bone marrow.

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development. Similarly, polypeptides and antibodies directed to these polypeptides are may be routinely detected in certain tissues and cell types (e.g., placenta, melanocytes, useful in providing immunological probes for differential identification of the tissue(s) iver, spleen, and bone marrow, and cancerous and wounded tissues) or bodily fluids sample taken from an individual having such a disorder, relative to the standard gene or cell type(s). For a number of disorders of the above tissues or cells, lower levels (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell expression level, i.e., the expression level in healthy tissue or bodily fluid from an Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: abnormal early individual not having the disorder.

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corresponding to this gene are useful for the treatment and diagnosis of abnormal early The tissue distribution indicates that polynucleotides and polypeptides development phenomena and diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 144 23

This gene is expressed primarily in fetal liver and spleen.

biological sample and for diagnosis of diseases and conditions: anemia and neutropenia. levels may be routinely detected in certain tissues and cell types (e.g., liver and spleen, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual providing immunological probes for differential identification of the tissue(s) or cell immune and blood systems, expression of this gene at significantly higher or lower type(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful as Similarly, polypeptides and antibodies directed to these polypeptides are useful in and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, reagents for differential identification of the tissue(s) or cell type(s) present in a

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having such a disorder, relative to the standard gene expression level, i.e., the

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expression level in healthy tissue or bodily fluid from an individual not thaving the

corresponding to this gene are useful in hematopoeisis and bone marrow regeneration as it is most abundant in fetal tissues responsible for the generation of hematopoeitic The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 145

lyrosine phosphatase which is thought to be important in transducing signal to activate The translation product of this gene shares sequence homology with protein cells such as T cell, B cell and other cell types.

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This gene is expressed primarily in T cells and tissues in early stages of development and to a lesser extent in cancers.

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: immuno-related reagents for differential identification of the tissue(s) or cell type(s) present in a diseases and cancer. Similarly, polypeptides and antibodies directed to these

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another tissue or cell sample taken from an individual having such a disorder, relative to ower levels may be routinely detected in certain tissues and cell types (e.g., embryonic the standard gene expression level, i.e., the expression level in healthy tissue or bodily tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or and fetal tissue, undifferentiated cells, and blood cells, and cancerous and wounded particularly of the immune system expression of this gene at significantly higher or 2 25

The tissue distribution and homology to the protein tyrosine phosphatase family indicate that polynucleotides and polypeptides corresponding to this gene are useful for luid from an individual not having the disorder. modulating the immune system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 146

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This gene is expressed primarily in T cell and to a lesser extent in B cell, macrophages and tumor tissues.

biological sample and for diagnosis of diseases and conditions: immuno disorders. Therefore, polynucleotides and polypeptides of the invention are useful as Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a

wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, routinely detected in certain tissues and cell types (e.g., blood cells, and cancerous and immune system, expression of this gene at significantly higher or lower levels may be type(s). For a number of disorders of the above tissues or cells, particularly of the providing immunological probes for differential identification of the tissue(s) or cel

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5 corresponding to this gene are useful for regulating the immune system therefore can be used in treating diseases such as autoimmune diseases and cancers. The tissue distribution indicates that polynucleotides and polypeptides

FEATURES OF PROTEIN ENCODED BY GENE NO: 147

cells, testis tumor, ovarian cancer, uterine cancer. This gene is expressed primarily in placenta and to a lesser extent in endothelial

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not limited to cancer. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions which include, but are reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

8 of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification endothelial cells, testis and ovary and other reproductive tissue, and cancerous and particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta,

25 wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue

corresponding to this gene are useful for diagnosis and treatment of cancers The tissue distribution indicates that polynucleotides and polypeptide:

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FEATURES OF PROTEIN ENCODED BY GENE NO: 148

Genet. 17, 40-48 (1997).) group cloned the human Torsin A gene. (See, Accession No. 2358279; see also Nature This sequence has significant homology to mouse torsin A. Recently, another

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standard gene expression level, i.e., the expression level in healthy tissue or bodily

fluid from an individual not having the disorder.

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lesser extent in fetal lung, fetal liver, fetal brain, adult brain and tumor tissues This gene is expressed primarily in osteoclastoma, T-cell, and placenta and to a

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5 and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, cells, bone, placenta, lung, liver, and brain and other tissues of the nervous system, or lower levels may be routinely detected in certain tissues and cell types (e.g., blood particularly of the hematopoiesis system, expression of this gene at significantly higher of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification hematopoiesis and cancers. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: disease conditions in expression level in healthy tissue or bodily fluid from an individual not having the having such a disorder, relative to the standard gene expression level, i.e., the synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

deficiencies in red blood cell, white blood cell, platelet and other hematopoiesis cells. corresponding to this gene are useful for treating blood related diseases such as The tissue distribution indicates that polynucleotides and polypeptides

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FEATURES OF PROTEIN ENCODED BY GENE NO: 149

20 spleen and osteoclastoma. endothelial cells and to a lesser extent in monocyte, dendritic cell, bone marrow, salivary gland, colon cancer, stomach cancer, pancreatic tumor, uterine cancer, fetal This gene is expressed primarily in T cell, prostate and prostate cancer,

25 မွ reagents for differential identification of the tissue(s) or cell type(s) present in a particularly of the immune system, expression of this gene at significantly higher or of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, diseases and cancers. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: immuno-related stomach, pancreas, uterus, spleen and bone, and cancerous and wounded tissues) or cells, prostate, endothelial cells, dendritic cells, bone marrow, salivary gland, colon, polypeptides are useful in providing immunological probes for differential identification tissue or cell sample taken from an individual having such a disorder, relative to the bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another lower levels may be routinely detected in certain tissues and cell types (e.g., blood Therefore, polynucleotides and polypeptides of the invention are useful as

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of cancers.

FEATURES OF PROTEIN ENCODED BY GENE NO: 150

This gene was recently cloned by another group, calling it eIF3-p66. (See Accession No. 2351378.) This gene plays a role in RNA binding and macromolecular assembly, and therefore, any mutations in this gene would likely result in a diseased phenotype. Preferred polypeptide fragments comprise the amino acid sequence: MAKFMTPVIQDNPSGWGPCAVPEQFRDMPYQPFSKGDRLGRVADWTGATYQDKRYTINKYSS

QFGGGSQYAYFHEEDESSFQLVDTARTQKTAYQRNRMRFAQRNLRRDKDRRNMLQFNLQLP
KSAKQKERERILQKKFQKQFGVRQKWDQKSQKPRDSSVEVRSDWEVKEEMDFPQLMKMRY
LEVSEPQDIECCGALEYYDKAFDRITTRSEKPLRXXKRIFHTVTTTDDPVIRKLAKTQGNVFATD
AILATLMSCTRSVYSWDIVVQRVGSKLFFDKRDNSDFDLLTVSETANEPPQDEGNSFNSPRNL
AMEATYINHNFSQQCLRMGKERYNFPNPFVEDDMDKNEIASVAYRYRSGKLGDDIDLIVRC

15 EHDGVMTGANGEVSFINIKTLNEWDSRHCNGVDWRQKLDSQRGAVIATELKNNSYKLARWTC
CALLAGSEYLKLGYVSRYHVKDSSRHVILGTQQFKPNEFASQINLSVENAWGILRCVIDICMKL
EEGKYLLIKDPNKQVIRVYSLPDGTFSS (SEQ ID NO: 638), as well as N-terminal and Cterminal deletions of this polypeptide fragment.

This gene is expressed primarily in T cell, bone marrow, embryo and endothelial cells and to a lesser extent in testis tumor and endometrial tumor.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune diseases and tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune disorders and cancers.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 151

This gene is expressed primarily in testis and to a lesser extent in T cell, spinal cord, placenta, neutrophil and monocyte.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: male reproductive and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testis and other reproductive tissue, blood cells, tissue of the nervous system, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating immune and reproductive functions.

FEATURES OF PROTEIN ENCODED BY GENE NO: 152

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The translation product of this gene shares sequence homology with tyrosyl-tRNA synthetase which is thought to be important in cell growth.

This gene is expressed primarily in brain, liver, keratinocytes, tonsils, and

25 heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer autoimmune diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, liver, keratinocytes, tonsils, heart expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissues of the nervous system, liver, keratinocytes, tonsils and heart, and cancerous and wounded tissues) or bodily

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fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or

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cell sample taken from an individual having such a disorder, relative to the standard

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gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

polynucleotides and polypeptides corresponding to this gene are useful for modulating The tissue distribution and homology to tyrosyl-tRNA synthetase indicate that

FEATURES OF PROTEIN ENCODED BY GENE NO:

Accession No. 2511745.) Dre4 is a gene required for steroidogenesis in Drosophila This gene is homologous to the Drosophila transcriptional regulator dre4. (See

5 5 melanogaster and encodes a developmentally expressed homologue of the yeast DPECIFFEQGGWSFL (SEQ ID NO: 639), as well as N-terminal and C-terminal deletions of FIEKVEALTKEELEFEVPFRDLGFNGAPYRSTCLLQPTSSALVNATEWPPFVVTLDEVELIHFXR acid sequence: KKRHTDVQFYTEVGEITTDLGKHQHMHDRDDLYAEQMEREMRHKLKTAFKN transcriptional regulator CDC68. Preferred polypeptide fragments comprise the amino this fragments. Also preferred are polynucleotide fragments encoding this polypeptide <u>VQFHLKNFDMVIVYKDYSKKVTMINAIPVASLDPIKEWLNSCDLKYTEGVQSLNWTKIMKTTVD</u>

This gene is expressed primarily in fetal liver, spleen, placenta, lung, T cell,

25 20 မွ or cell type(s). For a number of disorders of the above tissues or cells, particularly of plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from types (e.g., liver, spleen, placenta, lung, blood cells, thyroid, and testes and other significantly higher or lower levels may be routinely detected in certain tissues and cell the fetal liver, spleen, placenta, lung, T cell, thyroid, testes expression of this gene at useful in providing immunological probes for differential identification of the tissue(s) liver diseases. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: brain tumor, heart and reagents for differential identification of the tissue(s) or cell type(s) present in a the expression level in healthy tissue or bodily fluid from an individual not having the an individual having such a disorder, relative to the standard gene expression level, i.e., reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, Therefore, polynucleotides and polypeptides of the invention are useful as

FEATURES OF PROTEIN ENCODED BY GENE NO: 154

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testis, spleen, lung. This gene is expressed primarily in brain and to a lesser extent in fetal heart

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and other reproductive tissue, spleen, and lung, and cancerous and wounded tissues) or these polypeptides are useful in providing immunological probes for differential tissue or cell sample taken from an individual having such a disorder, relative to the bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissues and cell types (e.g., brain and other tissue of the nervous system, heart, testes this gene at significantly higher or lower levels may be routinely detected in certain tissues or cells, particularly of the brain, fetal heart, testis, spleen, lung expression of identification of the tissue(s) or cell type(s). For a number of disorders of the above diseases, immunological diseases. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: heart, liver and spleer reagents for differential identification of the tissue(s) or cell type(s) present in a fluid from an individual not having the disorder. standard gene expression level, i.e., the expression level in healthy tissue or bodily Therefore, polynucleolides and polypeptides of the invention are useful as

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FEATURES OF PROTEIN ENCODED BY GENE NO: 155

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25 that may provide a link between TCR activation and cell cycle control. essential to the completion of mitosis in yeast. Both endogenous and expressed murine proteins, and the mammalian homolog of Saccharomyces cerevisiae cdc48p, a protein protein (VCP). VCP is a member of a family of ATP binding, homo-oligomeric a 100 kDa protein. This gene is the human equivalent of murine valosin containing rapid tyrosine phosphorylation of a number of cellular proteins, one of the earliest being VCP are tyrosine phosphorylated in response to T cell activation. Thus we have identified a novel component of the TCR mediated tyrosine kinase activation pathway Activation of T cells through the T cell antigen receptor (TCR) results in the

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not limited to, cancer immunological disorders. Similarly, polypeptides and antibodies tissues and cell types (e.g., brain and other tissue of the nervous system, liver, spleen, differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, liver, spleen, placenta expression of directed to these polypeptides are useful in providing immunological probes for biological sample and for diagnosis of diseases and conditions which include, but are reagents for differential identification of the tissue(s) or cell type(s) present in a plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, this gene at significantly higher or lower levels may be routinely detected in certain Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in brain, liver, spleen, placenta

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an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to VCR indicate that polynucleotides and polypeptides corresponding to this gene are useful for treating cancer.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 156

The translation product of this gene shares sequence homology with rat growth response protein which is thought to be important in cell growth. A group recently cloned the human homolog of this gene, calling it insulin induced protein 1. (See Accession No. 2358269, see also, Genomics 43 (3), 278-284 (1997).) Preferred

10 cloned the human homolog of this gene, calling it insulin induced protein 1. (See Accession No. 2358269, see also, Genomics 43 (3), 278-284 (1997).) Preferred polypeptide fragments comprise the amino acid sequence: RSGLGLGHTAFLATLITQF LVYNGVYQYTSPDFLYIRSWLPCIFFSGGVTVGNIGRQLAMGVPEKPHSD (SEQ ID NO: 640). as well as N-terminal and C-terminal deletions of this polypeptide fragment. Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in brain, liver, placenta, heart, spleen,

lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for

differential identification of the tissue(s) or cell type(s). For a number of disorders of

the above tissues or cells, particularly of the brain, liver, placenta, heart, spleen.

expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, liver, placenta, heart, spleen, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to growth-response protein indicate that polynucleotides and polypeptides corresponding to this gene are useful for modulating cell growth.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 157

This gene is expressed primarily in Glioblastoma, endometrial tumor, lymphoma and pancreas tumor.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Glioblastoma, Endometrial tumor, lymphoma and pancreas tumor. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium, lymphoid tissue, pancreas, and tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the

FEATURES OF PROTEIN ENCODED BY GENE NO: 158

expression level in healthy tissue or bodily fluid from an individual not having the

The translation product of this gene shares sequence homology with IGE receptor which is thought to be important in allergy and asthma.

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This gene is expressed primarily in T cell, and fetal liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: allergy and asthma and other immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and

synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine,

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The tissue distribution and homology to IgE receptor indicate that polynucleotides and polypeptides corresponding to this gene are useful for allergy and asthma.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 159

The translation product of this gene shares sequence homology with immunoglobin heavy chain which is thought to be important in immune response to the antigen.

This gene is expressed primarily in activated neutrophil and to a lesser extent in activated T cell, monocyte and heart.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: infection, inflammation and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and heart, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinál fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to immunoglobin heavy chain variable region indicate that polynucleotides and polypeptides corresponding to this gene are useful for making the ligand to block specific antigen which cause certain disease.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 160

The translation product of this gene shares sequence homology with mouse X inactive specific transcript protein which is thought to be important in X chromosome

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This gene is expressed primarily in HSA172 cell and to a lesser extent in normal ovary tissue, ovarian cancer, frontal cortex and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

35 biological sample and for diagnosis of diseases and conditions: ovarian tumor, schizophrenia and other neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for

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differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and neural system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., ovary and other reproductive tissue, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to X inactive specific transcript protein indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of reproductive system tumors and CNS tumors.

FEATURES OF PROTEIN ENCODED BY GENE NO: 161

This gene is expressed primarily in adipose cell and to a lesser extent in liver and prostate.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: obesity and liver disorder. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the adipose cell, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adipose cells, liver, and

- prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- 30 The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of obesity and liver disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 162

The translation product of this gene shares sequence homology with yeast ubiquitin activating enzyme homolog which is thought to be important in protein posttraslation processing.

This gene is expressed primarily in stromal cell and to a lesser extent in retina, H. Atrophic Endometrium, colon carcinoma and myeloid progenitor cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of stromal cell development, neuronal growth disorders and tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and neural system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., retinal cells, endometrium, colon, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to ubiquitin-activating enzyme homolog indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of some type of tumors, fucosidosis and neuronal growth disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 163

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This gene is expressed primarily in primary breast cancer and nemangiopericytoma and to a lesser extent in adult brain and cerebellum.

polypeptides are useful in providing immunological probes for differential identification such a disorder, relative to the standard gene expression level, i.e., the expression level biological sample and for diagnosis of diseases and conditions: breast cancer, leukemia cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial significantly higher or lower levels may be routinely detected in certain tissues and cell of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, fluid or spinal fluid) or another tissue or cell sample taken from an individual having Therefore, polynucleotides and polypeptides of the invention are useful as and cerebellum disorders. Similarly, polypeptides and antibodies directed to these particularly of the immune system and neural system, expression of this gene at types (e.g., mammary tissue, brain and other tissue of the nervous system, and reagents for differential identification of the tissue(s) or cell type(s) present in a n healthy tissue or bodily fluid from an individual not having the disorder. 25 റ്റ 35

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of various tumors and disease involved in neural system.

5 FEATURES OF PROTEIN ENCODED BY GENE NO: 164

The translation product of this gene shares sequence homology with proline rich proteins. Recently, another group has also cloned this gene, calling it CD84 leukocyte antigen, a new member of the Ig superfamily. (See Accession No. U82988, see also, Blood 90 (6), 2398-2405 (1997).)

10 This gene is expressed primarily in Weizmann olfactory tissue and osteoclastoma and to a lesser extent in anergic T-cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: ostsis and immune 15 disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., olfactory tissue, bone, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution and homology to the Ig superfamily indicate that the protein product of this clone is useful for treatment of osteoporosis, autoimmune disease, and other immune disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 165

30 This gene is expressed primarily in atrophic endometrium and colon cancer and to a lesser extent in some fetal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a

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number of disorders of the above tissues or cells, particularly of the immune system,

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expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium, colon, and fetal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of tumors, specifically endometrium and colon tumors.

FEATURES OF PROTEIN ENCODED BY GENE NO: 166

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This gene is expressed primarily in human primary breast cancer and to a lesser extent in activated monocyte. Although the predicted signal sequence is identified in Table 1, other upstream sequences are also relevant. Preferred polypeptide fragments comprise the amino acid sequence: VTQPKHLSASMGGSVEIPFSFYYPWELAXXPXVRISWRRGHFHG QSFYSTRPPSIHKDYVNRLFLNWTEGQESGFLRISNLRKEDQSVYFCRVELDTRRSG (SEQ ID NO: 641), as well as N-terminal and C-terminal deletions. Also preferred are polypucleotide fragments encoding these polypeptide fragments.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: breast cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected

in certain tissues and cell types (e.g., mammary tissue, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of breast cancer.

FEATURES OF PROTEIN ENCODED BY GENE NO: 167

This gene is expressed primarily in fetal tissues and to a lesser extent in adult lung. This gene has also been mapped to chromosomal location 9q34, and thus, can be used as a marker for linkage analysis for chromosome 9.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or, cell type(s). For a number of disorders of the above tissues or cells, particularly of the embryo tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., fetal tissues, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e. the expression level in healthy tissue or bodily fluid from an individual not having the

FEATURES OF PROTEIN ENCODED BY GENE NO: 168

The translation product of this gene shares sequence homology with Ig Heavy Chain which is thought to be important in immune response.

This gene is expressed primarily in prostate cancer tissue specifically Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: prostate cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., prostate, tissue and cells of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the 30 disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 169

The translation product of this gene shares sequence homology with cytosolic acyl coenzyme-A hydrolase, which is thought to be important in neuron-specific fatty acid metabolism. The gene represented by this contig has since been published by Hajra and colleagues (GenBank Accession No. U91316).

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This gene is expressed primarily in human pituitary gland and to a lesser extent in colorectal cancer tissue. This gene has also been observed in the LNCAP cell line.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hyperlipidemias of familial and/or idiopathic origins. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly blood, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., pituitary and colon, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution and homology to rat cytosolic acyl coenzyme-A hydrolase indicate that polynucleotides and polypeptides corresponding to this gene are useful for the detection or treatment of hyperlipidemia disease states by virtue of the ability of specific drugs to activate the enzyme.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 170

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The translation product of this gene shares sequence homology with a Caenorhabditis elegans gene which is thought to be important in organism development.

This gene is expressed primarily in human synovial sarcoma tissue, bone marrow, and to a lesser extent in human brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, of bone, specifically synovial sarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the bone, connective tissues and possibly immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., synovial tissue, bone marrow, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another

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tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Caenorhabditis elegans indicate that polynucleotides and polypeptides corresponding to this gene are useful as a diagnostic and/or therapeutic modality directed at the detection and/or treatment of connective tissue sarcomas or other related bone diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 171

10 The translation product of this gene shares sequence homology with beta1-6GlcNAc transferase which is thought to be important in the transfer and metabolism of beta1-6, N-acetylglucosamine. This gene product has previously been shown to suppress melanoma lung metastasis in both syngeneic and nude mice, decreased invasiveness into the matrigel, and inhibition of cell attachment to collagen and laminin without affecting cell growth.

This gene is expressed primarily in human testes and prostate tissues, and to a lesser extent in kidney, medulla, and pancreas.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer particularly melanoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell

the above tissues or cells, particularly of the infiniture system, expression of this general significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testes and other reproductive tissue, prostate, kidney, pancreas, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to beta1-6GIcNAc transferase indicate that the protein product of this clone is useful for the development of diagnostic and/or therapeutic modalities directed at the detection and/or treatment of cancer, the metastasis of malignant tissue or cells. Defects in this potentially secreted enzyme may play a role in metastasis.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 172

This gene is expressed primarily in fetal spleen and liver.

5 5 S disorder, relative to the standard gene expression level, i.e., the expression level in routinely detected in certain tissues and cell types (e.g., spleen and liver, and cancerous immune systems, expression of this gene at significantly higher or lower levels may be number of disorders of the above tissues or cells, particularly of the hematopoiesis and polypeptides and antibodies directed to these polypeptides are useful in providing Wilm's tumor disease, hepatic disorders, and hematopoietic disorders. Similarly, biological sample and for diagnosis of diseases and conditions: immune disorders reagents for differential identification of the tissue(s) or cell type(s) present in a healthy tissue or bodily fluid from an individual not having the disorder. spinal fluid) or another tissue or cell sample taken from an individual having such a and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or immunological probes for differential identification of the tissue(s) or cell type(s). For Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and identification of fetal defects along with correcting diseases that affect hematopoiesis and the immune system.

FEATURES OF PROTEIN ENCODED BY GENE NO: 173

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The translation product of this gene shares sequence homology with ret II oncogene which is thought to be important in Hirschsprung disease and many types of cancers.

This gene is expressed in multiple tissues including the lymphatic system, brain, and thuroid

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Hirschsprung disease and multiple cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., lymphoid tissue, thyroid, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to

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the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ret II oncogene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of various cancers. It would also be useful for the diagnosis and treatment of Hirschsprung disease. Preferred polypeptides of the invention comprise the amino acid sequence: MEAQQVNEAESAREQLQXLHDQIAGQKASKQELETELERLKQEFHYIEEDLY RTKNTLQSRIKDRDEEIQKLRNQLTNKTLSNSSQSELENRLHQLTETLIQKQTMLESLSTEKNSL VFQLERLEQQMNSASGSSSNGSSINMSGIDNGEGTRLRNVPVLFNDTETNLAGMYGKVRKAAS OSDQFSIRLGIFLRRYPPIARVFVIIYMALLHLWVMIVLLTYTPEM HHDQPYGK (SEQ ID NO:

FEATURES OF PROTEIN ENCODED BY GENE NO: 174

The translation product of this gene shares sequence homology with testis enhanced gene transcript which is thought to be important in regulation of human development.

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This gene is expressed primarily in infant brain and to a lesser extent in a variety of other tissues and cell types, including the prostate, testes, monocytes, macrophages, dendritic cells, keratinocytes, and adipocytes.

20 30 25 of the above tissues or cells, particularly of the brain and immune systems, expression cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial of this gene at significantly higher or lower levels may be routinely detected in certain for differential identification of the tissue(s) or cell type(s). For a number of disorders antibodies directed to these polypeptides are useful in providing immunological probes developmental, immune and inflammation disorders. Similarly, polypeptides and biological sample and for diagnosis of diseases and conditions: neurological reagents for differential identification of the tissue(s) or cell type(s) present in a in healthy tissue or bodily fluid from an individual not having the disorder such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having testes and other reproductive tissue, blood cells, keratinocytes, and adipocytes, and tissues and cell types (e.g., brain and other tissue of the nervous system, prostate, Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution and homology to testis enhanced gene transcript indicate
that the protein product of this clone is useful for diagnosis and treatment of disorders involving the developing brain and the immune system.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 175

This gene is expressed primarily in prostate and to a lesser extent in various other tissues, including placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancers, especially of the prostate. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., prostate and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of prostate disorders and cancer. It may also be useful for the diagnosis and treatment of endocrine disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 176

The translation product of this gene shares sequence homology with Sacchromyces cerevisiae YNT20 gene which is thought to be important in mitochondrial function.

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This gene is expressed at a particularly high level in muscle tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases related to such tissues and cell types including: muscle wasting diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuromuscular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., muscle and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the YNT20 gene indicate that this protein is useful for treatment and detection of neuromuscular diseases caused by loss of mitochondrial function. For example this gene or its protein product could be used in replacement therapy for such diseases.

FEATURES OF PROTEIN ENCODED BY GENE NO: 177

This gene is expressed primarily in the brain and to a lesser extentin kidney, placenta, smooth muscle, heart and lung.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neuromuscular diseases, degenerative diseases of the central nervous system, and heart disease.

15 Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuromuscular system, central nervous system, and heart, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, kidney, placenta, muscle, heart and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

This gene or its protein product could also be used for replacement therapy for the above mentioned diseases.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 178

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The translation product of this gene shares sequence homology with caldesmon which is thought to be important in the cellular response to changes in glucose levels.

This gene is expressed primarily in multiple tissues including brain and retina.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: central nervous system disorders and retinopathy. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell

cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of disorders and retinopathy, expression of this gene at significantly higher or lower levels the nervous system, and retinal tissue, and cancerous and wounded tissues) or bodily type(s). For a number of disorders of the above tissues or cells, particularly of the CNS

and polypeptides corresponding to this gene are useful for treatment of retinopathies. The tissue distribution and homology to caldesmon indicate that polynucleotides

individual not having the disorder.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 179

fibrosin protein which is thought to be important in regulation of fibrinogenesis in The translation product of this gene shares sequence homology with mouse

15 certain chronic inflammatory diseases.

Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in amniotic cells and breast tissue

biological sample and for diagnosis of breast cancer and abnormal embryo reagents for differential identification of the tissue(s) or cell type(s) present in a

- 20 development. Similarly, polypeptides and antibodies directed to these polypeptides are or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) may be routinely detected in certain tissues and cell types (e.g., amniotic cells, and the reproductive system, expression of this gene at significantly higher or lower levels
- ß mammary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from the expression level in healthy tissue or bodily fluid from an individual not having the
- 30 of this gene is useful in the treatment of chronic inflammatory diseases could be used in replacement therapy for breast cancer. In addition the protein product of this clone is useful for treatment of breast cancer. This gene or its protein product The tissue distribution and homology to fibrosin indicate that the protein product

33 FEATURES OF PROTEIN ENCODED BY GENE NO: 180

various adult tissues including brain, lung, liver, testes, and prostate. This gene is expressed several infant tissues including brain and liver and

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5 biological sample and for diagnosis of diseases and conditions which include, but are system. Similarly, polypeptides and antibodies directed to these polypeptides are useful individual not having the disorder. gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an other reproductive tissue, and prostate, and cancerous and wounded tissues) or bodily and cell types (e.g., brain and other tissue of the nervous system, lung, liver, testes and gene at significantly higher or lower levels may be routinely detected in certain tissues central nervous system, hepatic system, and reproductive system, expression of this type(s). For a number of disorders of the above tissues or cells, particularly of the in providing immunological probes for differential identification of the tissue(s) or cell cell sample taken from an individual having such a disorder, relative to the standard fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or not limited to, brain cancer, lung cancer, liver cancer and cancers of the reproductive reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

growth blocker in a variety of settings including treatment of cancers. this clone is involved in growth regulation and could be used as a growth factor or The tissue distribution of this gene product indicates that the protein product of 5

20 FEATURES OF PROTEIN ENCODED BY GENE NO: 181

melanocytes and dendritic cells. This gene is expressed primarily in activated monocytes and to a lesser extent in

25 မွ in certain tissues and cell types (e.g., blood cells, melanocytes, and dendritic cells, and biological sample and for diagnosis of immune system diseases and cancer. Similarly, expression of this gene at significantly higher or lower levels may be routinely detected number of disorders of the above tissues or cells, particularly of the immune system, polypeptides and antibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial immunological probes for differential identification of the tissue(s) or cell type(s). For a fluid or spinal fluid) or another tissue or cell sample taken from an individual having Therefore, polynucleotides and polypeptides of the invention are useful as

ઝ in a variety of settings including treatment of cancers involved in growth regulation and could be used as a growth factor or growth blocker The tissue distribution indicates that the protein product of this clone could be

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This gene is expressed primarily in placenta and several tumors of various tissue origin and to a lesser extent in normal tissues including liver, lung, brain, and skin,

FEATURES OF PROTEIN ENCODED BY GENE NO: 182

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of cancers of all kinds. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, respiratory system and skin, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, lung, brain and other tissues of the nervous system, and skin, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The high expression of this gene in multiple tumors indicates that the protein product of the clone may be involved in cell growth control and therefore would be useful for treatment of certain cancers. Likewise molecules developed to block the activity of the protein product of this clone could be used to block its potential role in tumor growth promotion.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 183

The translation product of this gene shares sequence homology with the mouse Ndr1 gene which is thought to be important in cancer progression.

This gene is expressed multiple cell types and tissues including brain, lung, kidney, bone marrow, liver, and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as

or eagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of all types of cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous, immune, and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, lung, kidney, bone marrow, liver and spleen, and cancerous and wounded

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tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to Ndr1 gene, which is thought to be involved in cancer progression, indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of certain cancers. Likewise molecules developed to block the activity of the protein product of this clone could be used to block its potential role in tumor growth promotion.

FEATURES OF PROTEIN ENCODED BY GENE NO: 184

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This gene is expressed primarily in early stage human brain and liver and to a lesser extent in several other fetal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions; brain and liver cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, liver, and fetal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

The expression of this gene in embryonic tissues indicates that the protein could be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

or bodily fluid from an individual not having the disorder.

FEATURES OF PROTEIN ENCODED BY GENE NO: 185

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This gene is expressed primarily in infant and embryonic brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of degenerative nervous system disorders and brain cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The expression of this gene in embryonic tissues indicates that the protein could 10 be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

FEATURES OF PROTEIN ENCODED BY GENE NO: 186

This gene is expressed primarily in multiple tissues including placenta, fetal lung, fetal liver, and brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of all types of cancers including liver, brain and lung. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, pulmonary system, and hepatic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, lung, liver, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The expression of this gene in embryonic tissues indicates that the protein could be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZEQ SEQ SO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	•	SEQ NO: Y	First AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
1	HTTEZ21	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	11	582	1	582	177	177	313	1	18	19	22
1	HTTEZ21	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	197	1020	296	830	442	442	499	1	18	19	22
2	HBGBW52	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	12	465	1	465	81	81	314	1	30	31	128
2	HBGBW52	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	198	524	229	343		196	500	1	20	21	33
3	HCUFM41	97897 02/26/97 209043 05/15/97	ZAP Express	13	474	1	474	1	1	315		24	25	28
3	HCUFM41	97897 02/26/97 209043	ZAP Express	199	332	1	319	35	35	501	-1	24	25	28

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Gene	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	z g a g x	Total NT Seq.	of Clone Seg.		5' NT of Start Codon	•	SEQ DO: Y	AA of Sig Pep	Last AA of Sig Pep	Predicted First AA of Secreted Portion	
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	23	1486	596	1418	102	102	325	1			
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	203	847	1	839	87	87	505	ı	30	31	75
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	204	852	75	850		690	506	1			10
13	HTXEF04	209235 09/04/97	Uni-ZAP XR	205	1354	54	1354	100	100	507	1	33	34	207
14	HPMFD84	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	24	2323	1017	2059	1242	1242	326		21	22	68
14	HPMFD84	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	206	1378	113	1226	303	303	508		25	26	36
15	HE6DB26	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	25	683	1	683	304	304	327	1	30	31	84

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zges: xges:	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	•	SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ		AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
15	HE6DB26	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	207	1166	281	884	567	567	509	1	18	19	19
16	HHFFL33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	26	2036	14	1959	214	214	328	1	20	21	36
17	HODBD33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	27	717	1	717	70	70	329	1	30	31	63
17	HODBD33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	208	697	2	697	33	33	510	1	31	32	32
18	HMDAE90	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	28	495	1	495	39	39	330	1	24	25	35
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izeJ AA Io FIGO	AA iziiFi First AA fo forested notitoof	AA of Sig Pep	AA of gig 9-g	ASEQ BO NO: Y	to szri7 to AA Signal qs9	S' NT of Start Codon	of Clone Seq.	S' NT of Clone Seq.	Total NT Seq.	NO: REO SEO		ATCC Deposit No: S and Date	cDNA Clone ID	Сепе Йо.
	betribard	126 1	,33,13	_ ~ ~	IN S	L	TM 'F	TM '2		TN				

- -[·E		-oz-		-686-	-107-		-G7E-	1	-346-	-15-	-ЯХ-ЧАZ-inU-	26/51/50 10607 26/97/70 286846	-74ZUASH	_ LZ
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. It	18	30	τ	LEE	961	961	£\$6	ī	£†6	SE	XX 4AS-inU	L6/S1/S0 10007 16/97/70 86816	невеве6	57
84	ΙE	30.	Ī	988	\$8	58	LEV	I	L£ \$	34	pBluescript	L6/\$1/\$0 \$\$0607 L6/97/70 L68L6	H2KN115	7 7
izeJ AA io igo	Predicted AA triff o lo Screted hortion	AA Io	First A Of Sig Pep	≰ <mark></mark> ∯Aÿ≻	7' NT of sirst To AA Signal Pep	S; NT of Start Start Codon	of Clone Seq.	5' NT of Clone Seq.	Total NT Seq	Egabö×		ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgagx	Total NT Seq.	of	3' NT of Clone Seq.	Codon	AA of Signal Pep	SE E SE Y	AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
28	HSSDM73	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	38	672	1	672	22	22	340	1	38	39	42
29	HBMVK68	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	39	1908	135	1908	309	309	341	1	20	21	26
30	HMKDC66	97898 02/26/97 209044 05/15/97	pSport1	40	458	93	458	147	147	342	1	24	25	26
31	HMKCU94	97898 02/26/97 209044 05/15/97	pSport1	41	1153	500	1153	427	427	343	1	30	31	157
31	HMKCU94	97898 02/26/97 209044 05/15/97	pSport1	213	1079	502	896		739	515	1	23	24	43
32	HRDEW41	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	42	1983	1092	1983	27	27	344	1	11	12	520

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	FSES:x	Total NT Seq.	Seq.	of Clone Seq.	5' NT	5' NT of First AA of Signal Pep	SEQ	First AA of Sig Pep	AA of	Predicted First AA of Secreted Portion	Last AA of ORF
32	HRDEW41	97898 02/26/97 209044 05/15/97	Uni-ZAP XR				3357				1	10	20	39
33	HTOJN06	97898 02/26/97 209044 05/15/97	Uni-ZAP XR		1406	1	695		19	345	1	19	20	
34	HBGDA21	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	44	1391	851	1153	74	74	346	1	30	31	234
34	HBGDA21	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	215	1334	822	1036		638	517	1	18	19	174
35	HFGAK75		Uni-ZAP XR	45	1569	768	1569	14	14	347	1	19	20	169
35	HFGAK75	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	216	1511	770	1404	844	844	518		32	33	43

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εī			1	615	844		185	761	749	71Z	AX 4AS-inU	26/51/50 16/97/70 26/97/70	HBICW48	68
⊅ ∠Z	1£	30	ī	156	221	LLI	1300	788	9981	67	AX 4AS-inU	76/21/20 10607 16/97/70	HBICW¢8	68
\$7	70	61	ī	320	19	19	948	ī	346	817	AX 4AS-inU	05/17(6) 108074 108/37(6)		38
81	8£	4 £	ı	6 † E	Ibi	141	968	727	SLÞ	L Þ	pSport	L6/S1/S0 ++0607 L6/97/70 868L6	HOACE83	Lε
£\$	20	61	ı	348	79	79	1891	ī	1924		Uni-ZAP XR	76/21/20 10/27/20 16/92/20		9£
Last AA Io ISO	Predicted AA szri To Secreted Sortion		AA Jo BiZ	≰ ജ́вё≻	7, NT of First O AA Signal Signal	S' NT of Start Codon			Total NT Seq.	∑ Б≌ В Б В	Vector	ATCC Deposit No: X and Date	cDNA Clone ID	Gene No.

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ DO: NO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	Start Codon	5' NT of First AA of Signal Pep	SEQ NO: Y	AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
44	HMABL38	97899 02/26/97 209045 05/15/97	Uni-ZAP XR		1258	149	1190	254	254	522	1	18	19	26
45	HSKDK47	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	55	1896		1614	650	650	357	ı	33	34	47
46	HOSFH03	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	56	1753	555	1753	414	414	358	1	18	19	73
46	HOSFH03	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	221	1693	554	1693		526	523	1	25	26	58
47	HOGAV75	97899 02/26/97 209045 05/15/97	pCMVSport 2.0	57	1220	690	1024	128	128	359	I	30	31	102
47	HOGAV75	97899 02/26/97 209045 05/15/97	pCMVSport 2.0	222	1196	712	1163		1097	524	1			19

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	FSE BS: X	Total NT Seg.	Seq.	of Clone Seq.	5' NT of Start Codon	AA of Signal Pep	SEQ NO: Y		AA of Sig Pep		AA of ORF
48	HFCAI74	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	58	1049	362	1049	335	335	360	1	33	34	48
49	HAGBI17	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	59	1776	854	1737	189	189	361	1	30	31	179
49	HAGBI17	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	223	1791	979	1791	1164	1164	525	1	18	19	40
50	HLFBC91	97899 02/26/97 209045 05/15/97	pBluescript SK-	60	443	1	443	164	164	362	1	21	22	25
51	HPRCA31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	61	2888	1909	2888	90	90	363	1	30	31	224
51	HPRCA31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	224	2517	1597	2517	1953	1953	526	1	18	19	57

10 349 349	Secreted Portion 15	Sig 0£	Sig Pep I	725 7 7 7	Fep 139	Start Codon 139		565 1268	2424 1821 2ed: NL	\$22 29 X :ON	Yector RX 4AX-inU RX 4AX-inU	2:0N 2:0N 2:0N 2:0N 2:0N 2:0N 2:0N 2:0N 2:0N 2:0N 2:0N	HbKCE62 Cloue ID CDNY	Jene No. 52 52
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgas, x	Total NT Seq.	of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon		SEQ NO: Y	AA of Sig Pep	AA of Sig Pep		Last AA of ORF
59	HCWEF90	97899 02/26/97 209045 05/15/97	ZAP Express		448	9	448		1	532	1	22	23	75
60	HHGCM20	97899 02/26/97 209045 05/15/97	Lambda ZAP II	70	245	1	245	93	93	372	, ,	1	2	51
61	HFRAU10	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	71	361	1	361	1	1	373	1	30	31	61
61	HFRAU10	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	231	407	1	407	210	210	533	1	17	18	60
62	HATDT67	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	72	713	8	713	169	169	374	1	30	31	40
62	HATDT67	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	232	830	190	580	329	329	534	1	28	29	39

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgeb; x	NT Seq.	5' NT of Clone Seq.	of Clone Seq.	Start Codon	•	SEAS:>		AA of Sig Pep	Predicted First AA of Secreted Portion	
63	HOUBG93	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	73	862	1	862	67	67	375		30	31	
63	HOUBG93	97900 02/26/97 209046 05/15/97	Uni-ZAP XR		932	138	905	287	287	535	1			2
64	HMWEX24	97900 02/26/97 209046 05/15/97	Uni-Zap XR	74	4602		4525	730	730	376	1	30	31	203
64	HMWEX24	97900 02/26/97 209046 05/15/97	Uni-Zap XR	234	2786	2406	2739	2577	2577	536	1	22	23	36
65	HSGBA84	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	75	1255	1	1195	112	112	377	1	28	29	29
66	HTOCD52	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	76	475	1	475	13	13	378	1	30	31	136

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67	87	LZ	ī	382	767	767	5821	135	5821	08	AX 9AS-inU	L6/S1/S0 9 1 0607 L6/97/70 006L6	НОВИСЕ	04
67	17	20	1	185	<i>L</i> 97	<i>L</i> 97	8911	981	8911	6 <i>L</i>	AX 4AZ-inU	20807 9 1 0607 26/97/70		69
.81			ī	852	ISZ	152	111	ī	165	536	pBluescript	L6/\$1/\$0 9†0607 L6/97/70 006L6	нкіхве	89
894	18	30	1	380	56	97	0871	<i>L</i> 791	L 061	8 <i>L</i>	pBluescript	02/17/0 97/0607 26/97/70 006/6	нкіхве	89
ΙÞ	: † E	55	ī	6 <i>L</i> E	ħL	ΦL	667	52	\$9 1	LL	AX 4AZ-inU	26/51/50 9†0607 26/97/70 00626	914ЭЭДН	L9
ÞΙ			ī	LES	56	97	854	1	8S 1	527	AX 4AS-inU	26/51/50 9 1 0607 26/92/70 00626	HLOCDS2	99
izeal AA Io FISO	Predicted AA izri Po Secreted Rottion		First A of Sig Pep	≰ <mark>⊗</mark> Bö≻	S' NT of First AA of Signal Signal	S' NT of Start Codon	10	S' NT Of Clone Seq.	Total NT Seq.	ESO SECTION IN	101 39 V	ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.

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ı	to ARO	Secreted Portion	gi2 na9	SiS qs4	.V	Signal Pep	Start Codon	Seq.	Seq.	Seq.	X	Vector	No: X and Date	Clone ID	Gene. No.
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZSESX SEX	NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon		SEQ PO:Y	First AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
75	нвіавз9	97900 02/26/97 209046 05/15/97	Uni-ZAP XR		734	1	734	1	1	540	1	37	38	108
75	HBIAB39	97900 02/26/97 209046 05/15/97	Uni-ZAP XR		809	80	794		294	541	1	15	16	106
76	HTXDU73	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	86	1238	36	918	17	17	388	1			1
77	HOEAS24	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	87	1460	9	1458	166	166	389	1	53	54	299
77	HOEAS24	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	240	2201	841	2080	507	507	542	1	43	44	136
77	HOEAS24	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	241	1661	311	1520	390	390	543	1	35	36	424

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zges:x	Total NT Seq.	5' NT of Clone Seq.	of Clone Seg.	5' NT of Start Codon	•	Y	AA of Sig	AA of Sig Pep	Portion	Last AA of ORF
78	HTEIY30	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	88	1395	567	1395	639	639	390	1	36	37	49
79	HSKNE46	97900 02/26/97 209046 05/15/97	pBluescript	89	1186	352	1186	540	540	391	1	49	50	61
79	HSKNE46	97900 02/26/97 209046 05/15/97	pBluescript	242	1146	329	1146	564	564	544	1	21	22	39
80	HPMFL27	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	90	1821	1203	1614	1503	1503	392	1	30	31	79
81	HMWDN32	97900 02/26/97 209046 05/15/97		91	862	253	862	359	359	393	1	32	33	36
82	HPRAX55	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	92	696	349	696	98	98	394	1	30	31	180

19 £	77	17	ı	Lts	521	152	LESI	Ī	L ES1	542	AX 4AZ-inU	76/21/20 76/92/20 76/92/20		98
084	ΙE	0ε	ı	866	734	73 4	1082	811	1082	96	AX 4AZ-inU	76/21/20 76/92/20 76/92/20	HTPEG42	98
EE	ız	50	ı	945	161	161	116	7L	6751	***	AAS sbdms.I II	76/21/20 76/92/20 76/92/20	НСОРУЗЗ	\$8
ZS I	εε	32	1	<i>L</i> 6ε	907	907	8491	ī	2203	56	AX 4AZ-inU	20607 920607	НЗДЕЛ5	58
09	77	17	ı	968	\$8 <i>L</i>	58 <i>L</i>	ZLLI	ZVL	⊅ LLI	b 6	AX 4AZ-inU	L6/S1/S0 L7060Z L6/9Z/Z0 106L6	негыги	1 /8
17			ı	568	<i>L</i> 61	<i>L</i> 61	6541	1	9881	ε6	AX 4AS-inU	26/51/50 9 1 0607 26/97/70	ннггw36	٤8
85	εε	35	ī	545	348	848	0621	597	0561			02/12/61 9 1 /06/07 2/5/97/70		28
Last AA Io Igo	betoiteed AA tzri-T to Secreted notition	Lass A of Sig Pep	tzrif AA To Sig Gog	SEQ	S' NT of First o AA Signal Pep	5' NT of Start Codon	of Clone Seq.		Total NT Seq.	ZEÓ ZEÓ ZEÓ ZEÓ	10133V	ATCC Deposit No: Z and Date	cDNA Clone ID	jene Vo.

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ı		- 01	01	<u> </u>	100	234	731	2161	3/1	7171	- 60	dy d A S :-II	L6/\$1/\$0	Ozidaiki	- 00
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ı	ORE	Portion	Pep	Pep	I I	Pep	Codon			Seq.	l x	Vector	and Date	Clone ID	.oV
	10	Secreted	giZ	giZ	:ON	Signal	Start	Seq.	Seg.	IN	:ON		Z:0N	ANGo	Gene
ı	AA	30	30	Ιο	Œ	îo AA	ìo		Clone	Total	ID.		Deposit		
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ NO: N	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon		SEQ NO: Y		AA of Sig Pep		Last AA of ORF
91	HTSEL31	97901 02/26/97 209047 05/15/97	pBluescript	101	1394	608	1346	602	602	403	1	23	24	87
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	102	794	Î	794	518	518	404	1	30	31	92
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	248	1766	42	1766	356	356	550	1	30	31	168
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	249	2664	47	1708		147	551	1	18	19	124
93	HODAS59	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	103	1544	898	1531	975	975	405	1			21
94	НЕ6СТ48	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	104	871	106	871	248	248	406	1	34	35	174

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1 1				NT		5' NT	3' NT		of	AA	First	tast	Predicted	1
l I		ATCC		SEO		of	of	5' NT	First	SEQ			First AA	
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Gene	cDNA	No: Z		NO:	NT	Seq.	Seq.	Start	Signal		Sig	Sig		of
No.	Clone ID	and Date	Vector	X.	Seq.	ooq.	oug.	Codon		Y	Pep	Pep		ORF
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94	HE6CT48	97901	Uni-ZAP XR	250	865	97	865	258	258	552	1	19	20	177
1 1		02/26/97											ļ .	i 1
		209047					1	1		l	i	l	l '	l l
1		05/15/97					L		<u></u>			<u> </u>		
95	HMDAA61	97901	Uni-ZAP XR	105	404	1	404	16	16	407	1	21	22	64
1		02/26/97		l i		1	1	Ì	! .	ļ	1	1	ł	
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95	HMDAA61	97901	Uni-ZAP XR	251	2082	852	2074	829	829	553	1	22	23	72
1 1		02/26/97		1	i				l	Ī	l	1	1	
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		05/15/97					1	1		1	 	ļ.,.	 	000
96	HAQBK61	97901	Uni-ZAP XR	106	1542	506	1542	122	122	408	1	51	52	280
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		05/15/97		<u> </u>			<u> </u>		<u> </u>	l	ļ	<u> </u>	<u> </u>	
96	HAQBK61	97901	Uni-ZAP XR	252	1482	508	1482	ŀ	633	554	1	15	16	45
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96	HCUHB01	209215		253	834	1	834	82	82	555	1 I	40	41	251
		08/21/97		<u> </u>		<u> </u>			L	<u> </u>	<u> </u>	<u> </u>		L
97	HAQBF73	97901	Uni-ZAP XR	107	2327	1528	2327	465	465	409	1.1	30	31	284
	1	02/26/97		1	1	ı	1		1	1	Į	1	1	1
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761	LÞ	94	ι	717	Þ	Þ	15/1	696	1541	011	AAZ sbdms.J II	L6/S1/S0 L40607 L6/97/70 106L6	HLQAB52	
78	εī	71	ī	855	ISII		8877		LSEZ			L6/S1/S0 L 1 0607 L6/97/70 106L6	нетнеот	
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LEZ	15	30	<u> </u>	114	£06	9 <u>41</u>	<u>2431</u>	76S SLZ	7214		AX 4AZ-inU AX 4AZ-inU	106L6 L6/\$1/\$0 L 0 /97/70 106L6	HETHE07	
781 200	67	87		410	Z/1	7/1	7901	<u> </u>	7901	801	AX 4AZ-inU	10020 10020 10007 10020 10020		
61			,	955	886		8051	288	8051	552	AX 4A≤-inU	206751/20 70907 7097/20 70979		
Last AA of ORF	AA 12114 To Secreted noinof	to Sig gaq	AA Jo Sig Pep	X	First To AA Signal Pep	30	Seq.		Seq.	B No X	Vector	Deposit No: Z and Date	Clone ID	3
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1	801	57	74	I	195	242	747	1132	69	1193	526	Other	LZ960Z	HIBEK16	101
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	56	61	81	1	655	314	314	559	218	689	LSZ	AAZ abdmaJ	106L6	HLQAB52	001
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ DO: NO: X	Total NT Seq.	Seq.	of Clone Seg.	Start Codon		SEQ SEQ: Y	AA of Sig Pep	AA of Sig Pep		Last AA of ORF
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express			540	1171	337	337	416		30	31	163
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express	261		626	1161	335	335	563	1	30	31	253
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express	262	1162	629	1131	942	942	564	1			18
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	115	842	373	800	100	100	417	1	65	66	174
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	263	735	290	735			565	1			
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	264	783	416	783		413	566	1	33	34	73

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ DO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEQ ID		AA of	Predicted First AA of Secreted Portion 31	Last AA of ORF
106	НТЗАМ65	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	116		187	1470	581			1	30	31	263
106	НТЗАМ65	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	265	1638	301	1405	119	119	567				
106	НТЗАМ65	97901 02/26/97 209047 05/15/97	1	266	1455	148	1188	438	438	568		24	25	70
107	HE6DK18	97901 02/26/97 209047 05/15/97			952	418	906	499	499	419		28	29	120
108	НЕВЕК93	97901 02/26/97 209047 05/15/97		118	1256		1079		301	420		30	31	159
108	НЕВЕК93	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	267	1086	25	1050	227	227	569	1	23	24	34

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. 504	33	35	ı	777	881	8£1	07/1	ī	78 <i>L</i> 1	150	AX 4AZ-inU	L6/S1/S0 L70607 L6/97/70	_	011
721	_ 87	LZ	<u>I</u>	1 <i>L</i> S	737	737	5101	ÞΔI	1534	697	AX 4AZ-inU		нурсміо	601
101	32	34	1	0 <u>/</u> 5	SII	\$11	1003	17	1003	897	AX 4AZ-inU	26/51/50 26/97/70 26/97/70	НЉСМІО	601
₽ \$1	īs	ΟS	I	124	SLI	SLI	1501	1/1	£#11	611	AX 4AZ-inU	L6/S1/S0 L6/97/Z0 L6/97/Z0 106L6	ньсміо	601
128.1 АА 10 ЭЯО	Predicted AA izriF To Secreted noinor		AA 10 3i2	XO: REO SEO	S' NT of First To AA Oignal Signal	S' NT of Start Codon	of Clone Seq.	S' NT of Clone Seq.	Total TN Seq.	XEQ NO: NO: X		ATCC Deposit No: Z And Date	cDNA Clone ID	Gene No.

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82	61	81	1	ELS	988	988	1979	688	1571	172	AX 4AZ-inU	106L6	HEAAR60	EII
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zges:	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon		SEQ ID NO: Y		AA of Sig Pep	Portion	Last AA of ORF
116	HGBGK76	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		431	1	431	73	73	428	1	38	39	47
116	HGBGK76	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		515	1	515	43	43	-575	1	20	21	30
117	HBMUW78	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		3752	3465	3752	748	748	429	ľ	30	31	370
117	HBMUW78	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		2995	2738	2995	2777	2777	576		18	19	29
118	HASAS24	97902 02/26/97 209048 05/15/97		128	1144	669	1144	896	896	430	1			30
119	HSIDN55	97902 02/26/97 209048 05/15/97	·	129	1830	1234	1830	1265	1265	431	1			24

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	FSESX SESS	Total NT Seq.	of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon		SEQ:	First AA of Sig Pep	AA of Sig Pep	Portion	Last AA of ORF
120	HGBGZ64	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		1864	1505	1741	1578	1578	432		37	38	53
121	Н6ЕВЈ64	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	131	2041	1	1214	46	46	433	1	35	36	176
121	Н6ЕВЈ64	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	275	1990	8	1128	71	71	577	1	16	17	92
122	HOECP43	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	132	2012	853	1986	1127	1127	434	1	22	23	77
123	H2CBV31	97902 02/26/97 209048 05/15/97	pBluescript SK-	133		670	1632		962	435	1	25	26	32
124	HPCAD23	97902 02/26/97 209048 05/15/97		134	1565	281	1565	274	274	436	I	25	26	30

493						,									
PCT/US98/04493	ες	97	52	1	ltt	282	585	1347	ZLS	9440	681	AX 4AS-inU	20626 20607 20626 20626	ннгслег	156
*	33	61	81	τ	440	£811	£811	⊅6 ∠1	1044	SE61	138	Inoq2q	20626 20607 20626 20626	S6UAYIH	128
i	7 7	18	30	ı	872	97 <i>L</i>	97 <i>L</i>	2436	ZLS	2436	912	9AS sbdmsJ II	20626 20607 20626 20626	8+ннспн	LZI
	£†	15	30	1	654	184	†81	9061	τ	9061	151	AAS abdma.J II	20675 2090 2090 20972 2097	8+ннѕон	LZ1
	61			ī	438	401	401	0811	ī	1671		XX ¶AΣ-i∩U	L6/S1/S0 81/0607 L6/97/70	негонзі	156
	69	40	68	1	754		1124		1011			Inoged	26/\$1/\$0 8†0607 26/92/70 20626	SIDAGSH	125
	AA of TRO		lo gi2 qsq	io gi2 qsq	X	Io AA Ignai Qs¶	Содоп	Seq.		Total NT Seq.	NO:	Vector	Deposit No: S and Date	cDNA Clone ID	Gene No.
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	320	97	52	I	544	SS	ŞŞ	1595	ŞŞ	1569	143	4AZ sbdms.1	70616	H2G2C60	££1
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							İ						20607 26/97/20		
	0L	35	31	I	645	ILS		187	408	Z8 <i>L</i>	LLZ	AX 4AS-i⊓U	70676	HEBFU93	135
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	68	31	30	I	444	11	I	697	ī	697	745	AX 4AZ-inU	20676	невьпоз	135
													<i>L6/51/50</i>		ŀ
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•	34	·		I_	6443	\$6.	<u></u> ξ6_	L6 †	6	L6\$	141	AX 4AZ-inU	70676	HEBGA37	131
٠			l '							ŀ			L6/51/50	1	
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	7 9	52	74	I	245	9 <i>L</i> 9	919	6011	689	6011	140	AX 4AZ-inU	20676	LYTAD57	130
	ORE	Роплоп	Pep	Pep	J	dəd	Coqou			Seq.	х	Vector	and Date	Clone ID	.oN
	30	Secreted	Sig	gi2	:ON	Isngi 2		Seq.		JN	ON		Z:0N	ANGo	Gene
1	.AA	30	30	30	ID.	ìo AA	30		Slone	[BIOT]	ID,		Deposit		ŀ
	126.J	Predicted AA 121i7		tzni7 AA	SEÓ	io iziiT	TN 'S	1 NT C	TN 'S Îo		2EÓ NL		DOTA		ļ. I
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ NO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	•	SEQ NO: Y	First AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
135	HPTVC60	97902 02/26/97 209048 05/15/97	pBluescript	145	1021	526	1021	74	74	447	1	30	31	278
135	HPTVC60	97902 02/26/97 209048 05/15/97	pBluescript	278	961	524	961	545	545	580	1	23	24	110
136	HSKNE18	97902 02/26/97 209048 05/15/97	pBluescript	146	1285	5	1285	116	116	448	1	30	31	199
136	HSKNE18	97902 02/26/97 209048 05/15/97	pBluescript	279	1228	- 9	1228	324	324	581	1	26	27	30
137	HMWIF35	97902 02/26/97 209048 05/15/97	Uni-Zap XR	147	1386	169	1272	165	165	449	1	30	31	258
137	HMWIF35	97902 02/26/97 209048 05/15/97	Uni-Zap XR	280	1327	169	1208	160	160	582	1	23	24	71

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ ID NO: X	Total NT Seq.	5' NT of Clone Seq.	of	5' NT of Start Codon	AA of Signal	SEQ ID	AA of Sig	AA of Sig	Predicted First AA of Secreted Portion	
138	HMWGI25	97902 02/26/97 209048 05/15/97	Uni-Zap XR											
139	HSKGF03	97902 02/26/97 209048 05/15/97	pBluescript	149	1847	1689	1847	241	241	451]	33	34	315
139	HSKGF03	97902 02/26/97 209048 05/15/97	pBluescript	281	799	1	799		243	583	1	12	13	47
140	HMSKE75	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	150	1569	113	1517	417	417	452	1	21	22	52
141	HCMSH30	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	151	1540	538	1540	48	48	453	1	30	31	383
141	HCMSH30	97902 02/26/97 209048 05/15/97		282	2196	270	2196	294	294	584		32	33	39

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OL.	77	23	\perp	857	1 202	1592	6081	8741	1861	951	AX AAS-inII	FUOLO	LINVNSH	9/1
овъ	Portion	Pep.	Pep	X.	Pep	Содоп			Seq	х	Vector	and Date	Clone ID	.oV
jo	Secreted	Sig	Big	:ON	Signal	Start	Seq.	Seq.	TN	:ON		Z:ON	CDNA	эпэĐ
٧٧ I	10	10	jo	ď	10 AA	10		Clone	Total	Œ,		Deposit		
rası	AA izii I	٧٧	₩	SEQ	าราเป	TN 'S	10	Jo.		SEO		DOTA		
- 1	Predicted		First	AA	10	2.14 13		TN 'S	•	TN				
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70	77	71	1	954	017	07	212	I	1011	751	AX 4AS-inU	706L6	EIDMA'TH	144
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61	1	l	lτ	585	179	179	9911	LLZ	5811	283	psiuescript	706L6	HBMDW46	143
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163	LZ	97	lτ	557	561	561	٤98	1	£98	123	bgrnescubr	20676	HBMDM46	143
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981	53	25	1	†S †	9	9	SLST	069	6171	125	Inoq2q	70676	HLMCB92	745
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126.J	AA mii		AA	SEO	iziH	TN 'S	10	10		SEO		DDTA .	i	1
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L6/E0/L0
6E160Z
L6/S1/S0
6Y060Z
L6/9Z/Z0
6Y060Z
L6/9Z/Z0
6Y060Z
L6/9Z/Z0
E06L6
L6/S1/S0
6Y060Z
L6/9Z/Z0
E06L6
L6/S1/S0
6Y060Z
L6/9Z/Z0
E06L6
L6/S1/S0
6Y060Z
L6/9Z/Z0
E06L6

HCQAV96

H2KCO76

HS1AP03

HCFBC03

HCFBC03

HZNYKI

H2NYKI.

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II

AAZ abdmaJ

pBluescript

AX 9AZ-inU

pSport

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgesx	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	Start Codon	5' NT of First AA of Signal Pep	SEQ ID NO: Y	AA of Sig Pep		Predicted First AA of Secreted Portion	Last AA of ORF
150	HSHCC16	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		2120	1223		1416	1416		1			14
151	HTLEF62	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		900	482	900	46	46	463	1	30	31	285
151	HTLEF62	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		1517	783	1517	1062	1062	590]			24
152	HTLAD94	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	162	1003	1	1003	288	288	464	1	30	31	80
152	HTLAD94	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		3865	217	1195		281	591	1	16	17	38
153	HTSFQ12	97903 02/26/97 209049 05/15/97	pBluescript	163	2196	1607	2180	1611	1611	465	1	30	31	47

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	5805×	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	<u> </u>	SEQ DO: NO: Y		AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
154	HE6FL83	97903 02/26/97 209049 05/15/97		164	1945	271	1840	299	299	466	1	63	64	96
154	HE6FL83	97903 02/26/97 209049 05/15/97		290	1910	279	1818	355	355	592		39	40	69
155	HTXFJ55	97903 02/26/97 209049 05/15/97		165	2933	489	2871	258	258	467	1	30	31	399
155	HTXFJ55	97903 02/26/97 209049 05/15/97	İ	291	3276	486	2838		525	593	1	45	46	308
156	НЈРСЈ76	97903 02/26/97 209049 05/15/97	[166	2243	343	2221		341	468	1			1
157	HLTED27	97903 02/26/97 209049 05/15/97	Ì	167	1816	1130	1816	284	284	469	1	31	32	273

2															
2000001	35	54	53	ī	£/4	27 <i>L</i> 1	ZZL1	2100	7 1 91	2100	141	bBluescript	26/51/50 670607 26/97/70 20626	82ABWAH	191
2	74			ī	S65	015	015	1051	438	1051	£67	AX 4AS-inU	676750 670607 6797770 67903	HCECBSI	190
	102	91⁄2	SÞ	ī	7 <i>L</i> Þ	1001	1001	6981	86 <i>L</i>	£881			61/21/50 61/26/51 76/92/20	HCELB21	190
157	734	LZ	56	ī	14	_61	61	918	917	706	691	pBluescript	L6/51/50 61⁄0607 L6/97/70 20616	HNFIP24	651
22	761	61	81	I	04	802	802	L8L	Į	S † 6	891	Inoq2q	20303 203046 203770 20315 20315 20315 20315 20315 20315 20315	HWKBV¢4	851
	77			ı	76 5	1309	1306	8 1 51	8601	\$691	767	AX 4AS-iņU	L6/\$1/\$0 670607 L6/97/70 806L6	нстергл	۷\$1
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	ise.J	bətəibər¶ AA tzri刊		ızıi∓ AA	AA OBS	TN 'S Îo ÎsziF	TN 'S	TN 'E	TN '2		NT SEQ))		

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5' NT3' NT of of Clone Clone

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1630

Total NT Seq.

2892 762

567

<u>-828- 396 -828-</u>

Uni-ZAP XR

2K-bgjnescubt

2K-bBlaescubt

2K-bBInescubt

2K-bglnescubt

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Last AA Io Sig Pep

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208046 508046 L6/97/70

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163

193

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Gene No.

E06L6

L6/S1/S0
6+0607
L6/97/70
E06L6
L6/S1/S0
6+0607
L6/97/70
E06L6
L6/S1/S0
6+0607
L6/97/70
E06L6
L6/S1/S0
6+0607
L6/97/70
E06L6
L6/S1/S0
6+0607
L6/97/70
E06L6
L6/97/70
E06L6

ATCC Deposit No: Z and Date

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Last AA 10

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ NO: X	Total NT Seq.	of Clone Seq.		Start Codon		SEQ ID NO: Y	AA of Sig Pep	AA of Sig Pep	Portion	Last AA of ORF
165	HEAAL31	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		991	374	970	60	60	477	1	24	25	178
165	HEAAL31	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	297	2416	1387	2413	1473	1473	599	1	18	19	25
166	HFKFX55	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	176	1290	499	1290		688	478	1	25	26	52
167	H2LAO11	97903 02/26/97 209049 05/15/97	pBluescript SK-	177	2290	1	2290	173	173	479	1	22	23	62
168	HPFDZ95	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	178	549	1	549	11	11	480	1	21	22	27
168	HPFDZ95	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	298	545	1	545	17	17	600	1	21	22	27

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zeges: xeges:	Total NT Seq.	of	3' NT of Clone Seq.	5' NT of	5' NT of First AA of Signal Pep	SEQ ID	First AA of Sig Pep	AA of Sig	Predicted First AA of Secreted Portion	
169	HPTTUII	97904 02/26/97 209050 05/15/97	Uni-ZAP XR		1509	294	1352	92	92	481	1	30	31	339
169	HPTTUII	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	299	1530	385	1530	562	562	601	1	23	24	61
170	HCFAE79	97904 02/26/97 209050 05/15/97	pSport1	180	1316	985	1250	995	995	482	1	26	27	32
171	HTEDJ34	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	181	777	ì	777	51	51	483	1	30	31	48
171	HTEDJ34	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	300	997	244	997	300	300	602	1	23	24	29
172	HODCW06	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	182	791	1	791	14	14	484		29	30	38

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002	15	30	1	884	LSZ	LSZ	0811	794	7171	981	AX 9AS-inU	26/51/50 050607 26/97/70	1	941
54	01	6	I	109	09		9461	7	6987	302		L6/S1/S0 0S060Z L6/9Z/Z0		SLI
752	15	30	I	<u> </u>	<u> 19</u>	<i>L</i> 9	8877	SSE	2562	581	ЯХ ЧАХ-iaU	26/51/50 050607 26/97/70 20626	HE8BI92	SLI
69	LS	95	I	£09	533	233	2345	SL	2345	301	2K- bBluescript	26/51/50 050607 26/92/70 20626	H2MBF44	<i>\$L</i> 1
9 1 €	52	. 77	I	987	151	131	9651	\$L	9651	184	2K- bpjnescubt	L6/S1/S0 0S0607 L6/9Z/Z0 #06L6	H2MBF44	† †/I
19	12	07	ī	58⊅	SLS	SLS	5041	346	1402	£81	AX 4AZ-inU	26/\$1/\$0 050607 26/97/70	HFTAR26	£L1
izeal AA Io FISO	Predicted AA szrif Jo Secreted noriton	Last AA of Sig Pep	First AA Sig g∍9	≰ <mark>⊗</mark> Bö⊁	7. NT of first AA of Signal Pep	S' NT of Start Codon	of Clone Seq.	S' NT Clone Seq.	Total NT Seq.	Z S B S ×	Vector	ATCC Deposit No: Z and Date	cDNA Clone ID	Эспе Ио.

23							_						•		
PCT/US98/04493	_57_				-160-	-104-		-189-	-787-	-189-	-681-	-AX-4AS-inU	26/51/50 050607 26/97/70 50626	_rsaadah_	-6L1-
Đ.	52	61	81	I	ل	ъs	ьs	1561	I	£6 7 1		AX 4AZ-inU	6/51/50 050607 26/97/70		841
	597	15	<u>30</u>	ī	061	8	8	9151	096	9151		AX 9AS-inU	76/21/20 050602 76/92/20 10676		8/1
162	130	bb	£\$	1	909	L8L		sisi	61 <i>L</i>	7 £21	304	XX 4AZ-inU		HE6CWet	LLI
-	158	18_	30	11	68t	199	991	1224	0LL	1605	181	AX 4AS-inU		HE9CWet	<i>LL</i> I
	32	54	23	ī	\$09	£99	£99	6711	454	1811	303	AX 4AS-inU	76/51/50 26/97/70 76/97/70 706/6	HFTBR48	941
	1285.] AA 10 1900.	AA izriT 10 bataraa2		AA 10 gi2	KO: REO REO	triff To AA Signal qsq	S' NT of Start Codon	Clone Seq.		Total NT Seq.	SEQ ZOX	Vector	ATCC Deposit No: X and Date	cDNA Clone ID	Gene No.
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				NT			3, ML		of First		First AA		Predicted First AA	
		ATCC		SEQ ID.		of	of Clone	5' NT	AA of		of	of	of	AA
_	~~.	Deposit		NO:	Total NT			Start	Signal		Sig	Sig	Secreted	of
Gene	cDNA	No: Z		X		Seq.	Seq.	Codon		Y Y	Sig	Sig		ORF
No.	Cione ID	and Date	Vector	^	Seq.						Pep	Pep		
180	HCEEK08	97904	Uni-ZAP XR	190	1014	703	1014	360	360	492	1	30	31	159
		02/26/97								i	1	l .	1	1
ì	ĺ	209050								i	l	l	į	l
		05/15/97		202		<u> </u>			1,25	700	<u> </u>			
180	HCEEK08	97904	Uni-ZAP XR	306	577	1	577	l	175	608	1		1	6
	l	02/26/97			1	i	['	ł	İ	i ·	l	l	1 .	
	1	209050		1	i	1	1				1	1	ļ	1
	********	05/15/97	Discourant	101	2779	2207	2630	1153	1153	493	-	30	31	279
181	HAFAU18	97904	pBluescript SK-	191	2//9	2207	2030	1133	1133	493	'	30	31	2/9
		02/26/97 209050	21/2	1	ľ	l	l			ľ		1	l	ļ
	i	05/15/97			l		ì				i	l	1	
181	HAFAU18	97904	pBluescript	307	2860	163	2860	21	21	609	1	30	31	232
101	HAPAUIO	02/26/97	SK-	1307	2000	1 .05	12000	- 1.	1	1007	1 1	"	"	
	İ	209050	JIL.	ł			İ	l	l	1	ı			
	1	05/15/97			İ	1		j	1				1	
181	HAFAU18	97904	pBluescript	308	876	275	876	302	302	610	1	32	33	34
		02/26/97	SK-	l	İ	!		İ		l	ı	ł	Ī	l
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		05/15/97		<u> </u>			1		<u> </u>		<u> </u>			L
182	HETBY74	97904	Uni-ZAP XR	192	1923	30	1923	45	45	494	1	33	34	193
		02/26/97	ĺ	1	ł		1		1		1		Į.	1
	1	209050			l	l	ŀ		1	1		l	1	1
	1	05/15/97	I	I	ŀ	I	1	i	1	l	ŀ	1	i	i

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NEQ SEQ SO: NO: X	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	AA of Signal	SEQ ID		AA of	Predicted First AA of Secreted Portion	
183	HTOAF35	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	193		1160		178	178	495	1	30	31	205
183	HTOAF35	97904 02/26/97 209050 05/15/97		309	2025	840	2025	971	971	611	1	18	19	21
184	HCRBX32	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	194	3054	2004	3054	434	434	496	1	11	12	147
184	HCRBX32	97904 02/26/97 209050 05/15/97			3026	1966	3026		2131	612	1			9
185	HEBGB80	97904 02/26/97 209050 05/15/97			907	152	907	297	297	497	1	30	31	64
185	HEBGB80	97904 02/26/97 209050 05/15/97		311	712	67	712	107	107	613		18	19	29

ORE

128.J AA 10

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Portion

Secreted beloiber9 AA lzri7 lo Last AA of Sig Pep AA AA Io giZ gəq 86t

NO: ZEO VY

572

5' NT of First Signal Pep

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'cDNA clone ID" identified in Table 1 and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous sequence assembled from partially homologous ("overlapping") sequences obtained from the of high redundancy (usually three to five overlapping sequences at each nucleotide described above. The nucleotide sequence identified as "NT SEQ ID NO.X" was Table 1 summarizes the information corresponding to each "Gene No. position), resulting in a final sequence identified as SEQ ID NO:X.

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deposit number listed in "ATCC Deposit No:Z and Date." Some of the deposits contain multiple different clones corresponding to the same gene. "Vector" refers to the type of The cDNA Clone ID was deposited on the date and given the corresponding vector contained in the cDNA Clone ID. 2

the nucleotide position of SEQ ID NO: X of the predicted signal sequence is identified as Total NT Seq." refers to the total number of nucleotides in the contig identified reflected by the nucleotide position indicated as "5" NT of Clone Seq." and the "3" NT putative start codon (methionine) is identified as "5" NT of Start Codon." Similarly, of Clone Seq." of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the by "Gene No." The deposited clone may contain all or most of these sequences, "5' NT of First AA of Signal Pep." 15

LZ6

572

5° NT of Start Codon

687

608

3' NT of Clone Seq.

S8L

†8

of Clone Seq.

LN .9

6871

290

Total NT Seq.

961

X ID SEO

Vector

The translated amino acid sequence, beginning with the methionine, is identified alternative open reading frames are specifically contemplated by the present invention. as "AA SEQ ID NO:Y," although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by these 2

The first and last amino acid position of SEQ ID NO: Y of the predicted signal

peptide is identified as "First AA of Sig Pep" and "Last AA of Sig Pep." The predicted "Predicted First AA of Secreted Portion." Finally, the amino acid position of SEQ ID NO:Y of the last amino acid in the open reading frame is identified as "Last AA of first amino acid position of SEQ ID NO: Y of the secreted portion is identified ORF." 25

Deposit No: Z and Date

DOTA

981

981

Gene No.

HFAMH74

Clone ID

SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and

below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO; X or the cDNA be used to generate antibodies which bind specifically to the secreted proteins encoded molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO: Y may otherwise suitable for a variety of uses well known in the art and described further contained in the deposited clone. These probes will also hybridize to nucleic acid by the cDNA clones identified in Table 1. 8 35

sequence. In these cases, the predicted amino acid sequence diverges from the actual 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion amino acid sequence, even though the generated DNA sequence may be greater than sequencing errors. The errors exist as misidentified nucleotides, or as insertions or in an open reading frame of over 1000 bases) deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid deletions of nucleotides in the generated DNA sequence. The erroneously inserted or Nevertheless, DNA sequences generated by sequencing reactions can contain

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20 sequence or the amino acid sequence, the present invention provides not only the sequencing the deposited clone in accordance with known methods. The predicted amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA amino acid sequence can then be verified from such deposits. Moreover, the amino containing a human cDNA of the invention deposited with the ATCC, as set forth in generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated containing the deposited human cDNA, collecting the protein, and determining its determined by peptide sequencing or by expressing the protein in a suitable host cell acid sequence of the protein encoded by a particular clone can also be directly Table 1. The nucleotide sequence of each deposited clone can readily be determined by Accordingly, for those applications requiring precision in the nucleotide

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identifying or amplifying the corresponding gene from appropriate sources of genomic Such methods include preparing probes or primers from the disclosed sequence and accordance with known methods using the sequence information disclosed herein. SEQ ID NO: Y, or the deposited clone. The corresponding gene can be isolated in The present invention also relates to the genes corresponding to SEQ ID NO:X.

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sequences provided herein and screening a suitable nucleic acid source for the desired homologs may be isolated and identified by making suitable probes or primers from the Also provided in the present invention are species homologs. Species

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polypeptides, synthetically produced polypeptides, or polypeptides produced by a polypeptides include isolated naturally occurring polypeptides, recombinantly produced combination of these methods. Means for preparing such polypeptides are wel The polypeptides of the invention can be prepared in any suitable manner. Such

mature form, or may be a part of a larger protein, such as a fusion protein (see below). The polypeptides may be in the form of the secreted protein, including the ઝ

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such as multiple histidine residues, or an additional sequence for stability during secretory or leader sequences, pro-sequences, sequences which aid in purification, recombinant production. It is often advantageous to include an additional amino acid sequence which contains

5 using antibodies of the invention raised against the secreted protein in methods which one-step method described in Smith and Johnson, Gene 67:31-40 (1988) polypeptide, including the secreted polypeptide, can be substantially purified by the form, and preferably are substantially purified. A recombinantly produced version of a are well known in the art Polypeptides of the invention also can be purified from natural or recombinant sources The polypeptides of the present invention are preferably provided in an isolated

Signal Sequences

20 5 cleavage point for that sequence, are available. For instance, the method of McGeoch indicates the amino terminus of the secreted protein. The accuracy of predicting the produce the same predicted cleavage point(s) for a given protein. the range of 75-80%. (von Heinje, supra.) However, the two methods do not always cleavage points of known mammalian secretory proteins for each of these methods is in from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information region and a subsequent uncharged region of the complete (uncleaved) protein. The Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged Methods for predicting whether a protein has a signal sequence, as well as the

30 shown in Table 1. methods of McGeoch and von Heinje are incorporated. The analysis of the amino acid Engineering 10:1-6 (1997)), which predicts the cellular location of a protein based on was analyzed by a computer program called SignalP (Henrik Nielsen et al., Protein sequences of the secreted proteins described herein by this program provided the results the amino acid sequence. As part of this computational prediction of localization, the In the present case, the deduced amino acid sequence of the secreted polypeptide

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or - 5 residues) of the predicted cleavage point. Similarly, it is also recognized that in shown in SEQ ID NO:Y which have an N-terminus beginning within 5 residues (i.e., + Accordingly, the present invention provides secreted polypeptides having a sequence vary from organism to organism and cannot be predicted with absolute certainty. As one of ordinary skill would appreciate, however, cleavage sites sometimes

ડ્ડ some cases, cleavage of the signal sequence from a secreted protein is not entirely

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polynucleotides encoding such polypeptides, are contemplated by the present invention. uniform, resulting in more than one secreted species. These polypeptides, and the

secreted protein to the ER. These polypeptides, and the polynucleotides encoding such occurring signal sequence may be further upstream from the predicted signal sequence. necessarily predict the naturally occurring signal sequence. For example, the naturally However, it is likely that the predicted signal sequence will be capable of directing the Moreover, the signal sequence identified by the above analysis may not polypeptides, are contemplated by the present invention.

Polynucleotide and Polypeptide Variants

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polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical "Variant" refers to a polynucleotide or polypeptide differing from the to the polynucleotide or polypeptide of the present invention.

Applied Math 48:1073 (1988).) Methods commonly employed to determine identity or methods to measure identity between two polynucleotide or polypeptide sequences, the term "identity" is well known to skilled artisans. (Carillo, H., and Lipton, D., SIAM J Press, (1987); and SEQUENCE ANALYSIS PRIMER, Gribskov, M. and Devereux, published techniques. (See, e.g.: (COMPUTATIONAL MOLECULAR BIOLOGY, SEQUENCE ANALYSIS IN MOLECULAR BIOLOGY, von Heinje, G., Academic Research (1984) 12(1):387 (1984)), BLASTP, BLASTN, FASTA (Atschul, S.F. et INFORMATICS AND GENOME PROJECTS, Smith, D.W., ed., Academic Press, programs, including the GCG program package (Devereux, J., et al., Nucleic Acids Lesk, A.M., ed., Oxford University Press, New York, (1988); BIOCOMPUTING: "Identity" per se has an art-recognized meaning and can be calculated using similarity between two sequences include, but are not limited to, those disclosed in "Guide to Huge Computers," Martin J. Bishop, ed., Academic Press, San Diego, New York, (1993); COMPUTER ANALYSIS OF SEQUENCE DATA, PART I, i, eds., M Stockton Press, New York, (1991).) While there exists a number of (1994), and Carillo, H., and Lipton, D., SIAM J Applied Math 48:1073 (1988). Methods for aligning polynucleotides or polypeptides are codified in computer Griffin, A.M., and Griffin, H.G., eds., Humana Press, New Jersey, (1994); 15 20 25 8

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the reference polynucleotide and that gaps in identity of up to 5% of the total number of parameters are set so that the percentage of identity is calculated over the full length of When using any of the sequence alignment programs to determine whether a particular sequence is, for instance, 95% identical to a reference sequence, the nucleotides in the reference polynucleotide are allowed.

sequences or both amino acid sequences. The result of said global sequence alignment Penalty=1, Joining Penalty=30, Randomization Group Length=0, and Cutoff Score=1, sequence (a sequence of the present invention) and a subject sequence, also referred to calculate percent identity and similarity of an amino acid alignment are: Matrix=PAM length in nucleotide bases, whichever is shorter. Preferred parameters employed to (1990).) The term "sequence" includes nucleotide and amino acid sequences. In a Gap Penalty=5, Gap Size Penalty 0.05, and Window Size=500 or query sequence program based on the algorithm of Brutlag et al. (Comp. App. Biosci. 6:237-245 150, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group as a global sequence alignment, can be determined using the FASTDB computer A preferred method for determing the best overall match between a query is in percent identity. Preferred parameters used in a FASTDB search of a DNA sequence to calculate percent identiy are: Matrix=Unitary, k-tuple=4, Mismatch sequence alignment the query and subject sequences are either both nucleotide 9 2

sequence at least 95% identical to SEQ ID NO:X or the deposited clone, up to 5% of the As an illustration, a polynucleotide having a nucleotide sequence of at least 95% ive point mutations per each 100 nucleotides of the total length (not just within a given 00 nucleotide stretch). In other words, to obtain a polynucleotide having a nucleotide SEQ ID NO:X or the cDNA except that the polynucleotide sequence may include up to deposited clone, means that the polynucleotide is identical to a sequence contained in inserted, or substituted with other nucleotides. These changes may occur anywhere nucleotides in the sequence contained in SEQ ID NO:X or the cDNA can be deleted 'identity" to a sequence contained in SEQ ID NO:X or the cDNA contained in the hroughout the polynucleotide.

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Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty=0.05, and Window

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Size=500 or query sequence length in amino acid residues, whichever is shorter.

95%, 96%, 97%, 98% or 99% identity to a sequence contained in SEQ ID NO:X or the cDNA contained in the deposited clone. Of course, due to the degeneracy of the genetic Further embodiments of the present invention include polynucleotides having at the polynucleotides having at least 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity code, one of ordinary skill in the art will immediately recognize that a large number of east 85% identity, more preferably at least 90% identity, and most preferably at least

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575 Science Drive, Madison, WI 53711 (using the local homology algorithm of Smith

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and Waterman, Advances in Applied Mathematics 2:482-489 (1981).)

Package, Version 8 for Unix, Genetics Computer Group, University Research Park,

al., J. Molec. Biol. 215:403 (1990), Bestfit program (Wisconsin Sequence Analysis

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NO:Y or the expressed protein produced by the deposited clone will encode a polypeptide identical to an amino acid sequence contained in SEQ ID

sequence of the polypeptide is identical to the reference polypeptide except that the deleted or substituted with another amino acid, or a number of amino acids up to 5% of acids of the total length of the reference polypeptide. In other words, to obtain a polypeptide sequence may include up to five amino acid alterations per each 100 amino reference sequence or in one or more contiguous groups within the reference sequence carboxy terminal positions of the reference amino acid sequence or anywhere between sequence. These alterations of the reference sequence may occur at the amino or the total amino acid residues in the reference sequence may be inserted into the reference acid sequence, up to 5% of the amino acid residues in the reference sequence may be polypeptide having an amino acid sequence at least 95% identical to a reference amino example, 95% "identity" to a reference polypeptide, is intended that the amino acid those terminal positions, interspersed either individually among residues in the Similarly, by a polypeptide having an amino acid sequence having at least, for

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least 80% identity, more preferably at least 85% identity, more preferably at least 90% biological activity of the protein. the deposited clone. Preferably, the above polypeptides should exhibit at least one amino acid sequence contained in SEQ ID NO: Y or the expressed protein produced by identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to an Further embodiments of the present invention include polypeptides having at

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still more preferably at least 96%, 97%, 98%, or 99% similarity to an amino acid deposited clone. sequence contained in SEQ ID NO:Y or the expressed protein produced by the polypeptides having at least 90% similarity, more preferably at least 95% similarity, and In a preferred embodiment, polypeptides of the present invention include

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or both. Especially preferred are polynucleotide variants containing alterations which substitutions due to the degeneracy of the genetic code are preferred. Moreover, produce silent substitutions, additions, or deletions, but do not alter the properties or the human mRNA to those preferred by a bacterial host such as E. coli). combination are also preferred. Polynucleotide variants can be produced for a variety activities of the encoded polypeptide. Nucleotide variants produced by silent of reasons, e.g., to optimize codon expression for a particular host (change codons in variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any The variants may contain alterations in the coding regions, non-coding regions.

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several alternate forms of a gene occupying a given locus on a chromosome of an Naturally occurring variants are called "allelic variants," and refer to one of

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techniques or by direct synthesis. Alternatively, non-naturally occurring variants may be produced by mutagenesis allelic variants can vary at either the polynucleotide and/or polypeptide level organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).)

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carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).) exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma (1993), reported variant KGF proteins having heparin binding activity even after loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 deleted from the N-terminus or C-terminus of the secreted protein without substantial polypeptides of the present invention. For instance, one or more amino acids can be technology, variants may be generated to improve or alter the characteristics of the Using known methods of protein engineering and recombinant DNA

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8 2 amino acid position. The investigators found that "[m]ost of the molecule could be the entire length of the molecule. Multiple mutations were examined at every possible 3,500 individual IL-la mutants that averaged 2.5 amino acid changes per variant over analysis of human cytokine IL-1a. They used random mutagenesis to generate over coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational activity similar to that of the naturally occurring protein. For example, Gayle and sequences examined, produced a protein that significantly differed in activity from wildaltered with little effect on either [binding or biological activity]." (See, Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide Moreover, ample evidence demonstrates that variants often retain a biological

30 are removed from the N-terminus or C-terminus. Whether a particular polypeptide C-terminus of a polypeptide results in modification or loss of one or more biological will likely be retained when less than the majority of the residues of the secreted form deletion variant to induce and/or to bind antibodies which recognize the secreted form functions, other biological activities may still be retained. For example, the ability of a readily be determined by routine methods described herein and otherwise known in the lacking N- or C-terminal residues of a protein retains such immunogenic activities can Furthermore, even if deleting one or more amino acids from the N-terminus or

35 have little effect on activity. For example, guidance concerning how to make substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as Thus, the invention further includes polypeptide variants which show

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phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

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The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

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As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Glu, replacement of the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

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Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

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For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

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Polynucleotide and Polypeptide Fragments

In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in the deposited clone or shown in SEQ ID NO:X. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in the deposited clone or the nucleotide sequence shown in SEQ ID NO:X. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, 500, 600, 2000 nucleotides) are preferred.

Moreover, representative examples of polynucleotide fragments of the invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, and 701 to the end of SEQ ID NO:X or the cDNA contained in the deposited clone. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity.

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In the present invention, a "polypeptide fragment" refers to a short arnino acid sequence contained in SEQ ID NO.Y or encoded by the cDNA contained in the deposited clone. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, and 161 to the end of the coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90,

100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about"

includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

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Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

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Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity

Epitopes & Antibodies

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In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA

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81:3998-4002 (1983).)

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Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier. However,

immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is

meant to include intact molecules as well as antibody fragments (such as, for example,
Fab and F(ab')2 fragments) which are capable of specifically binding to protein. Fab
and F(ab')2 fragments lack the Fc fragment of intact antibody, clear more rapidly from
the circulation, and may have less non-specific tissue binding than an intact antibody.

(Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred
as well as the products of a FAB or other immunoglobulin expression library.

Moreover, antibodies of the present invention include chimeric, single chain, and
humanized antibodies.

Fusion Proteins

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Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

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Examples of domains that can be fused to polypeptides of the present invention

include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through

linker sequences.

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polypeptide to improve stability and persistence during purification from the host cell or preparation of the polypeptide. The addition of peptide moieties to facilitate handling of Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino polypeptide to facilitate purification. Such regions may be removed prior to final acids, particularly charged amino acids, may be added to the N-terminus of the subsequent handling and storage. Also, peptide moieties may be added to the polypeptides are familiar and routine techniques in the art.

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polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) facilitate purification and show an increased half-life in vivo. One reported example immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins Moreover, polypeptides of the present invention, including fragments, and describes chimeric proteins consisting of the first two domains of the human CD4monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. can also be more efficient in binding and neutralizing other molecules, than the specifically epitopes, can be combined with parts of the constant domain of Biochem. 270:3958-3964 (1995).)

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would be desired. For example, the Fc portion may hinder therapy and diagnosis if the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D. proteins comprising various portions of constant region of immunoglobulin molecules deleting the Fc part after the fusion protein has been expressed, detected, and purified, Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion together with another human protein or part thereof. In many cases, the Fc part in a Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. example, human proteins, such as hIL-5, have been fused with Fc portions for the example, improved pharmacokinctic properties. (EP-A 0232 262.) Alternatively, fusion protein is beneficial in therapy and diagnosis, and thus can result in, for fusion protein is used as an antigen for immunizations. In drug discovery, for Chem. 270:9459-9471 (1995).) 23

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sequences, such as a peptide which facilitates purification of the fused polypeptide. In Moreover, the polypeptides of the present invention can be fused to marker

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derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).) Chatsworth, CA, 91311), among others, many of which are commercially available. preferred embodiments, the marker amino acid sequence is a hexa-histidine pepude, Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue,

Thus, any of these above fusions can be engineered using the polynucieolides or the polypeptides of the claimed invention.

Vectors. Host Cells, and Protein Production

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The present invention also relates to vectors containing the polynucleotide of the vector. Retroviral vectors may be replication competent or replication defective. In the echniques. The vector may be, for example, a phage, plasmid, viral, or retroviral present invention, host cells, and the production of polypeptides by recombinant

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as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such a virus, it may be packaged in vitro using an appropriate packaging cell line and then latter case, viral propagation generally will occur only in complementing host cells.

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expression constructs will further contain sites for transcription initiation, termination, translation initiating codon at the beginning and a termination codon (UAA, UGA or promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to and, in the transcribed region, a ribosome binding site for translation. The coding name a few. Other suitable promoters will be known to the skilled artisan. The The polynucleotide insert should be operatively linked to an appropriate portion of the transcripts expressed by the constructs will preferably include a UAG) appropriately positioned at the end of the polypeptide to be translated. ransduced into host cells.

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resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance Streptomyces and Salmonella typhimurium cells; fungal cells, such as yeast cells; insect cells such as Drosophila S2 and Spodoptera Sf9 cells; animal cells such as CHO, COS, selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin As indicated, the expression vectors will preferably include at least one appropriate hosts include, but are not limited to, bacterial cells, such as E. coli, genes for culturing in E. coli and other bacteria. Representative examples of

293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and conditions for the above-described host cells are known in the art.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan.

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Introduction of the construct into the host cell can be effected by calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation, transduction, infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., Basic Methods In Molecular Biology (1986). It is specifically contemplated that the polypeptides of the present invention may in fact be expressed by a host cell lacking a recombinant vector.

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A polypeptide of this invention can be recovered and purified from recombinant cell cultures by well-known methods including armmonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for preferation.

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Polypeptides of the present invention, and preferably the secreted form, can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells. Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any protein after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

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Uses of the Polynucleotides

Each of the polynucleotides identified herein can be used in numerous ways as reagents. The following description should be considered exemplary and utilizes known techniques.

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The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat polymorphisms), are presently available. Each polynucleotide of the present invention can be used as a chromosome marker.

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Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be selected using computer analysis so that primers do not span more than one predicted exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment.

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Similarly, somatic hybrids provide a rapid method of PCR mapping the polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flowsorted chromosomes, and preselection by hybridization to construct chromosome

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specific-cDNA libraries.

Precise chromosomal location of the polynucleotides can also be achieved using
fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This
technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides
2,000-4,000 bp are preferred. For a review of this technique, see Verma et al.,
"Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York
(1988).

For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for marking multiple sites and/or multiple chromosomes). Preferred polynucleotides correspond to the noncoding regions of the cDNAs because the coding sequences are more likely conserved within gene families, thus increasing the chance of cross hybridization during chromosomal mapping.

Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage

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analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library).) Assuming I megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

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Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker.

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In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Both methods rely on binding of the polynucleotide to DNA or RNA. For these techniques, preferred polynucleotides are usually 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease.

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Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the

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present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell.

The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for

The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

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Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

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There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers specific to particular tissue prepared from the sequences of the present invention. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for

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In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

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Uses of the Polypeptides

Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypoptide of the present invention can be used to assay protein levels in a

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell. Biol. 105:3087-

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3096 (1987).) Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (1251, 1211), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technetium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and bickin

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In addition to assaying secreted protein levels in a biological sample, proteins can also be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

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A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 1311, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously, or intraperitoneally) into the mammal. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20

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millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. In vivo turnor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Turnor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982).)

Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an individual; (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

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Moreover, polypeptides of the present invention can be used to treat disease. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B), to inhibit the activity of a polypeptide (e.g., an oncogene), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth).

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Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

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At the very least, the polypeptides of the present invention could be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

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ological Activities

35 The polynucleotides and polypeptides of the present invention can be used in assays to test for one or more biological activities. If these polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules

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may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides could be used to treat the associated disease.

Immune Activity

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A polypeptide or polynucleotide of the present invention may be useful in treating deficiencies or disorders of the immune system, by activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune deficiencies or disorders may be genetic, somatic, such as cancer or some autoimmune disorders, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, a polynucleotide or polypeptide of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

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A polynucleotide or polypeptide of the present invention may be useful in treating or detecting deficiencies or disorders of hematopoietic cells. A polypeptide or polynucleotide of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat those disorders associated with a decrease in certain (or many) types hematopoietic cells. Examples of immunologic deficiency syndromes include, but are not limited to: blood protein disorders (e.g. agammaglobulinemia, dysgammaglobulinemia), ataxia telangiectasia, common variable immunodeficiency, Digeorge Syndrome, HIV infection, HTLV-BLV infection, leukocyte adhesion deficiency syndrome, lymphopenia, phagocyte bactericidal dysfunction, severe combined immunodeficiency (SCIDs), Wiskott-Aldrich Disorder, anemia, thrombocytopenia, or hemoglobinuria.

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Moreover, a polypeptide or polynucleotide of the present invention could also be used to modulate hemostatic (the stopping of bleeding) or thrombolytic activity (clot formation). For example, by increasing hemostatic or thrombolytic activity, a polynucleotide or polypeptide of the present invention could be used to treat blood coagulation disorders (e.g., afibrinogenemia, factor deficiencies), blood platelet disorders (e.g. thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, a polynucleotide or polypeptide of the present invention that can decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment of heart attacks (infarction), strokes, or scarring.

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A polynucleotide or polypeptide of the present invention may also be useful in treating or detecting autoimmune disorders. Many autoimmune disorders result from

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inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing

Examples of autoimmune disorders that can be treated or detected by the present invention include, but are not limited to: Addison's Disease, hemolytic anemia, antiphospholipid syndrome, rheumatoid arthritis, dermatitis, allergic encephalomyelitis, lo glomerulonephritis, Goodpasture's Syndrome, Graves' Disease, Multiple Sclerosis, Myasthenia Gravis, Neuritis, Ophthalmia, Bullous Pemphigoid, Pemphigus, Polyendocrinopathies, Purpura, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Thyroiditis, Systemic Lupus Erythematosus, Autoimmune Pulmonary Inflammation, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye disease.

Similarly, allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated by a polypeptide or polynucleotide of the present invention. Moreover, these molecules can be used to treat anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

A polynucleotide or polypeptide of the present invention may also be used to treat and/or prevent organ rejection or graft-versus-host disease (GVHD). Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. The administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T.

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an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing organ rejection or GVHD.

Similarly, a polypeptide or polynucleotide of the present invention may also be used to modulate inflammation. For example, the polypeptide or polynucleotide may inhibit the proliferation and differentiation of cells involved in an inflammatory response. These molecules can be used to treat inflammatory conditions, both chronic and acute conditions, including inflammation associated with infection (e.g., septic shock, sepsis, or systemic inflammatory response syndrome (SIRS)), ischemiateperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel

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disease, Crohn's disease, or resulting from over production of cytokines (e.g., TNF or

Hyperproliferative Disorders

A polypeptide or polynucleotide can be used to treat or detect hyperproliferative disorders, including neoplasms. A polypeptide or polynucleotide of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, a polypeptide or polynucleotide of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

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For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

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Examples of hyperproliferative disorders that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but are not limited to neoplasms located in the: abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

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Similarly, other hyperproliferative disorders can also be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome. Waldenstron's Macroplobulinemia. Gaucher's Disease, histiocytosis, and

25 Syndrome, Waldenstron's Macroglobulinemia, Gaucher's Disease, histiocytosis, and any other hyperproliferative disease, besides neoplasia, located in an organ system listed above.

Infectious Disease

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A polypeptide or polynucleotide of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, the polypeptide or polynucleotide of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

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Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus,

- 5 Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza), Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g.,
- Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiollitis, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), meningitis, opportunistic infections (e.g., AIDS),
- pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.
- 20 Similarly, bacterial or fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacterial families and fungi: Actinomycetales (e.g., Corynebacterium, Mycobacterium, Norcardia), Aspergillosis, Bacillaceae (e.g., Anthrax, Clostridium), Bacteroidaceae,
- Blastomycosis, Bordetella, Borrelia, Brucellosis, Candidiasis, Campylobacter, Coccidioidomycosis, Cryptococcosis, Dermatocycoses, Enterobacteriaceae (Klebsiella, Salmonella, Serratia, Yersinia), Erysipelothrix, Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales, Neisseriaceae (e.g., Acinetobacter, Gonorrhea, Menigococcal), Pasteurellacea Infections (e.g., Actinobacillus,
- Heamophilus, Pasteurella), Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, and Staphylococcal. These bacterial or fungal families can cause the following diseases or symptoms, including, but not limited to: bacteremia, endocarditis, eye infections (conjunctivitis, tuberculosis, uveitis), gingivitis, opportunistic infections (e.g., AIDS related infections), paronychia, prosthesis-related infections, Reiter's Disease, respiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, Dysentery, Paratyphoid Fever, food poisoning, Typhoid, pneumonia, Gonorrhea, meningitis, Chlamydia, Syphilis, Diphtheria,

impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections. A polypeptide or polynucleotide of the present invention can be used to treat or detect Leprosy, Paratuberculosis, Tuberculosis, Lupus, Botulism, gangrene, tetanus, any of these symptoms or diseases.

Malaria, pregnancy complications, and toxoplasmosis. A polypeptide or polynucleotide Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, These parasites can cause a variety of diseases or symptoms, including, but not limited Moreover, parasitic agents causing disease or symptoms that can be treated or giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), detected by a polynucleotide or polypeptide of the present invention include, but not Leishmaniasis, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas. of the present invention can be used to treat or detect any of these symptoms or to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, limited to, the following families: Amebiasis, Babesiosis, Coccidiosis,

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patient, or by removing cells from the patient, supplying the cells with a polynucleotide therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease. invention could either be by administering an effective amount of a polypeptide to the of the present invention, and returning the engineered cells to the patient (ex vivo Preferably, treatment using a polypeptide or polynucleotide of the present

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diseases.

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Regeneration

disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteocarthritis, periodontal Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, A polynucleotide or polypeptide of the present invention can be used to injury, or systemic cytokine damage. 25 39

skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac), vascular (including vascular endothelium), nervous, hematopoietic, and Tissues that could be regenerated using the present invention include organs without or decreased scarring. Regeneration also may include angiogenesis.

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Moreover, a polynucleotide or polypeptide of the present invention may increase avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel polypeptide of the present invention could also be used prophylactically in an effort to egeneration of non-healing wounds includes pressure ulcers, ulcers associated with regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. A polynucleotide or syndrome, and other tendon or ligament defects. A further example of tissue vascular insufficiency, surgical, and traumatic wounds.

Drager syndrome), could all be treated using the polynucleotide or polypeptide of the neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized polynucleotide or polypeptide of the present invention to proliferate and differentiate Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shystoke). Specifically, diseases associated with peripheral nerve injuries, peripheral disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Similarly, nerve and brain tissue could also be regenerated by using a present invention. 2 15

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A polynucleotide or polypeptide of the present invention may have chemotaxis fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial hyperproliferation. The mobilized cells can then fight off and/or heal the particular cells) to a particular site in the body, such as inflammation, infection, or site of activity. A chemotaxic molecule attracts or mobilizes cells (e.g., monocytes, trauma or abnormality.

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chemotaxic activity of particular cells. These chemotactic molecules can then be used to issues by attracting immune cells to the injured location. Chemotactic molecules of the disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotaxic molecules can be used to treat wounds and other trauma to treat inflammation, infection, hyperproliferative disorders, or any immune system present invention can also attract fibroblasts, which can be used to treat wounds. A polynucleotide or polypeptide of the present invention may increase

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invention may inhibit chemotactic activity. These molecules could also be used to treat It is also contemplated that a polynucleotide or polypeptide of the present

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disorders. Thus, a polynucleotide or polypeptide of the present invention could be used as an inhibitor of chemotaxis.

Binding Activity

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A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules

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Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

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Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from mammals, yeast, Drosophila, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the molecule to observe binding, stimulation, or inhibition of activity of either the

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The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

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polypeptide or the molecule.

Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule activity or binding, and comparing the polypeptide/molecule activity or binding to a standard

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Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The

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antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

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Other Activities

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A polypeptide or polynucleotide of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

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A polypeptide or polynucleotide of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide or polynucleotide of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

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A polypeptide or polynucleotide of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, caricadic rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity),

hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

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A polypeptide or polynucleotide of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional components.

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Other Preferred Embodiments

nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of Other preferred embodiments of the claimed invention include an isolated SEQ ID NO:X wherein X is any integer as defined in Table 1.

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positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of Clone Sequence and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Start Codon and ending with the nucleotide at about the position of the 3' Nucleotide of Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the Similarly preferred is a nucleic acid molecule wherein said sequence of range of positions beginning with the nucleotide at about the position of the 5' NO:X in Table 1

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sequence which is at least 95% identical to a sequence of at least about 150 contiguous Also preferred is an isolated nucleic acid molecule comprising a nucleotide nucleotides in the nucleotide sequence of SEQ ID NO:X.

sequence which is at least 95% identical to a sequence of at least about 500 contiguous Further preferred is an isolated nucleic acid molecule comprising a nucleotide nucleotides in the nucleotide sequence of SEQ ID NO:X.

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ID NO:X beginning with the nucleotide at about the position of the 5' Nucleotide of the nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in First Amino Acid of the Signal Peptide and ending with the nucleotide at about the A further preferred embodiment is a nucleic acid molecule comprising a

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A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NQ:X. 35

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to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or molecule which hybridizes does not hybridize under stringent hybridization conditions stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid Also preferred is an isolated nucleic acid molecule which hybridizes under of only T residues.

Also preferred is a composition of matter comprising a DNA molecule which which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1. for said comprises a human cDNA clone identified by a cDNA Clone Identifier in Table cDNA Clone Identifier. 2

Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the nucleotides in the nucleotide sequence of a human cDNA clone identified by a cDNA Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous ATCC Deposit Number shown in Table 1. Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said human cDNA clone.

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Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

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contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone. A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said human cDNA clone. 25

a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical A further preferred embodiment is a method for detecting in a biological sample group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer dentified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the notecule in said sample with a sequence selected from said group and determining to a sequence of at least 50 contiguous nucleotides in a sequence selected from the as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone comprises a step of comparing a nucleotide sequence of at least one nucleic acid ATCC Deposit Number shown for said cDNA clone in Table 1; which method 8 35

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whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

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A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

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Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

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Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1.

Also preferred is a polypeptide, wherein said sequence of contiguous amino acids is included in the amino acid sequence of SEQ ID NO: Y in the range of positions beginning with the residue at about the position of the First Amino Acid of the Secreted

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Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Frame as set forth for SEQ ID NO:Y in Table 1.

Portion and ending with the residue at about the Last Amino Acid of the Open Reading

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Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

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Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a secreted portion of the secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the

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clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with amino acid sequence of the secreted portion of the protein encoded by a human cDNA the ATCC Deposit Number shown for said cDNA clone in Table 1.

clone identified by a cDNA Clone Identifier in Table I and contained in the deposit with Also preferred is an isolated polypeptide comprising an amino acid sequence at amino acid sequence of the secreted portion of the protein encoded by a human cDNA least 95% identical to a sequence of at least about 100 contiguous amino acids in the the ATCC Deposit Number shown for said cDNA clone in Table 1.

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contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of the secreted portion of the protein 2

human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in sequence of at least 10 contiguous amino acids in a sequence selected from the group the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. consisting of: an amino acid sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a polypeptide comprising an arnino acid sequence that is at least 90% identical to a Further preferred is an isolated antibody which binds specifically to a 2

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least 10 contiguous amino acids in a sequence selected from the group consisting of: an Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the and a complete amino acid sequence of a protein encoded by a human cDNA clone molecule in said sample with a sequence selected from said group and determining comprises a step of comparing an amino acid sequence of at least one polypeptide whether the sequence of said polypeptide molecule in said sample is at least 90% ATCC Deposit Number shown for said cDNA clone in Table 1; which method identical to said sequence of at least 10 contiguous amino acids.

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comprising an amino acid sequence that is at least 90% identical to a sequence of at least Also preferred is the above method wherein said step of comparing an amino polypeptides in said sample to an antibody which binds specifically to a polypeptide 10 contiguous amino acids in a sequence selected from the group consisting of: an acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of 35

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amino acid sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1; identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the and a complete amino acid sequence of a protein encoded by a human cDNA clone ATCC Deposit Number shown for said cDNA clone in Table 1. Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. defined in Table 1; and a complete amino acid sequence of a secreted protein encoded Also preferred is a method for identifying the species, tissue or cell type of a consisting of: an amino acid sequence of SEQ ID NO: Y wherein Y is any integer as 2

Also preferred is the above method for identifying the species, tissue or cell type sequence of at least 10 contiguous amino acids in a sequence selected from the above sequences, wherein at least one sequence in said panel is at least 90% identical to a molecules comprising an amino acid sequence in a panel of at least two amino acid of a biological sample, which method comprises a step of detecting polypeptide

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obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two arnino acid sequences, wherein at least one sequence in said panel identified in Table 1, which method comprises a step of detecting in a biological sample Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number associated with abnormal structure or expression of a gene encoding a secreted protein Also preferred is a method for diagnosing in a subject a pathological condition sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid is at least 90% identical to a sequence of at least 10 contiguous amino acids in a shown for said cDNA clone in Table 1.

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In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

polypeptide wherein said polypeptide comprises an amino acid sequence that is at least Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding

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1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table any integer as defined in Table 1; and a complete amino acid sequence of a secreted from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected

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clone in Table 1.

in a prokaryotic host. sequence encoding a polypeptide has been optimized for expression of said polypeptide Also preferred is an isolated nucleic acid molecule, wherein said nucleotide

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2 ATCC Deposit Number shown for said cDNA clone in Table 1. identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the complete amino acid sequence of a secreted protein encoded by a human cDNA clone sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1; and a comprises an amino acid sequence selected from the group consisting of: an amino acid Also preferred is an isolated nucleic acid molecule, wherein said polypeptide

recombinant host cell produced by this method. a recombinant host cell comprising introducing the vector into a host cell, as well as the the recombinant vector produced by this method. Also preferred is a method of making inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is Further preferred is a method of making a recombinant vector comprising

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cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the and an amino acid sequence of a secreted portion of a protein encoded by a human of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y is defined in Table 1; NO:Y beginning with the residue at the position of the First Amino Acid of the Secreted sequence selected from the group consisting of: an amino acid sequence of SEQ ID polypeptide is a secreted portion of a human secreted protein comprising an amino acid expressed and recovering said polypeptide. Also preferred is this method of making an culturing this recombinant host cell under conditions such that said polypeptide is Portion of SEQ ID NO: Y wherein Y is an integer set forth in Table 1 and said position isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The Also preferred is a method of making an isolated polypeptide comprising

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individual a pharmaceutical composition comprising an amount of an isolated level of a secreted protein activity, which method comprises administering to such an Also preferred is a method of treatment of an individual in need of an increased

isolated polypeptide produced by this method is also preferred.

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the level of said protein activity in said individual polypeptide, polynucleotide, or antibody of the claimed invention effective to increase

understood by reference to the following examples, which are provided by way of Having generally described the invention, the same will be more readily

ķ illustration and are not intended as limiting.

Example 1: Isolation of a Selected cDNA Clone From the Deposited

5 15 example, where a particular clone is identified in Table 1 as being isolated in the vector was isolated. In many cases, the vector used to construct the library is a phage vector Table 1 identifies the vectors used to construct the cDNA library from which each clone related plasmid for each phage vector used in constructing the cDNA library. For from which a plasmid has been excised. The table immediately below correlates the Each cDNA clone in a cited ATCC deposit is contained in a plasmid vector.

"Lambda Zap," the corresponding deposited clone is in "pBluescript." Vector Used to Construct Library pBluescript (pBS) Corresponding Deposited Plasmid

pSport1 pCMVSport 3.0 pCMVSport 2.0 lafmid BA Zap Express pSport1 plafmid BA pBK pCMVSport 3.0 pCMVSport 2.0

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Uni-Zap XR Lambda Zap

pBluescript (pBS)

25 Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap pCR®2.1

5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., Nucleic Acids Res.

- ဗ contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1 Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS
- 35are the first sites on each respective end of the linker). "+" or "-" refer to the orientation sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which The S and K refers to the orientation of the polylinker to the T7 and T3 primer

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of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initiated from the f1 ori generates sense strand DNA and in the other, antisense.

Vectors pSport1, pCMVSport 2.0 and pCMVSport 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gnuber, C. E., et al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY) contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1 Blue. Vector pCR*2.1, which is available from Invitrogen, 1600 Faraday Avenue, Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed

10. Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark, J. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9: (1991).) Preferably, a polynucleotide of the present invention does not comprise the phage vector sequences identified for the particular clone in Table 1, as well as the corresponding plasmid vector sequences designated above.

The deposited material in the sample assigned the ATCC Deposit Number cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 50 plasmid DNAs, each containing a different cDNA clone; but such a deposit sample may include plasmids for more or less than 50 cDNA clones, up to about 500 cDNA clones.

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Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO.X.

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Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ¹²P-Y-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection

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agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1,93 to 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are

15 performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993).)

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Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

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This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged

prosputation in recessary to enrunate 3 prosputate groups on degraded or damaged

35 RNA which may interfere with the later RNA ligase step. The phosphatase should then
be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to

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remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be

ligated to an RNA oligonucleotide using T4 RNA ligase.

This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

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Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

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A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the cDNA sequence corresponding to SEQ ID NO:X., according to the method described in Example 1. (See also, Sambrook.)

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Example 3: Tissue Distribution of Polypeptide

Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method described in Example 1 is labeled with P³² using the rediprimeTM DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN-100TM column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for mRNA expression.

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Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) (Clontech) are examined with the labeled probe using ExpressHybTM hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are

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30 mounted and exposed to film at -70°C overnight, and the films developed according to standard procedures.

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(Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM

IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased

Example 4: Chromosomal Mapping of the Polynucleotides

An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of

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conditions: 30 seconds, 95°C; 1 minute, 56°C; 1 minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions is analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

Example 5: Bacterial Expression of a Polypeptide

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A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (AmpP), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

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The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (KanP). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis. Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D. 60) of between 0.4 and 0.6. IPTG

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Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic

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gene expression.

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QIAGEN, Inc., supra). Proteins with a 6 x His tag bind to the Ni-NTA resin with high onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from removed by centrifugation, and the supernatant containing the polypeptide is loaded agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., supra).

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Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

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The purified protein is then renatured by dialyzing it against phosphate-buffered recommended conditions are as follows: renature using a linear 6M-1M urea gradient in saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the renaturation the proteins are eluted by the addition of 250 mM immidazole. Immidazole 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer protein can be successfully refolded while immobilized on the Ni-NTA column. The The renaturation should be performed over a period of 1.5 hours or more. After plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

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In addition to the above expression vector, the present invention further includes a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgamo sequence, and 6) the lactose operon Number XXXXXX.) This vector contains: 1) a neomycinphosphotransferase gene as inked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession repressor gene (lacIq). The origin of replication (oriC) is derived from pUC19 (LTI, an expression vector comprising phage operator and promoter elements operatively Gaithersburg, MD). The promoter sequence and operator sequences are made synthetically. 2 23

Xbal, BamHI, Xhol, or Asp718, running the restricted product on a gel, and isolating nsert is generated according to the PCR protocol described in Example 1, using PCR the larger fragment (the stuffer fragment should be about 310 base pairs). The DNA DNA can be inserted into the pHEa by restricting the vector with Ndel and Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible primers having restriction sites for Ndel (5' primer) and Xbal, BamHI, Xhol, or enzymes. The insert and vector are ligated according to standard protocols. 3

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The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

Example 6: Purification of a Polypeptide from an Inclusion Body

The following alternative method can be used to purify a polypeptide expressed in E coli when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the $\it E.~coli$ fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at

- 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer. 2
- (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by The cells are then lysed by passing the solution through a microfluidizer NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4. 2
- 13 ndrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the The resulting washed inclusion bodies are solubilized with 1.5 M guanidine pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

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Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps. 22

filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a

stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0. The CM-20 column is then eluted using a 10 column

10 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0

The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from

15 Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded.

The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

Example 7: Cloning and Expression of a Polypeptide in a Baculovirus Expression System

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In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide into a baculovirus to express a polypeptide. This expression vector contains the strong polyhedrin promoter of the Autographa californica nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites such as BarnHI. Xba I and

- 25 Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from E. coli under control of a weak Drosophila promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated
- 30 homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

Many other baculovirus vectors can be used in place of the vector above, such as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription,

translation, secretion and the like, including a signal peptide and an in-frame AUG as

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required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-39 (1989).

Specifically, the cDNA sequence contained in the deposited clone, including the

AUG initiation codon and the naturally associated leader sequence identified in Table 1, is amplified using the PCR protocol described in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the pA2 vector does not need a second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures,"

10 Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.).

The fragment and the dephosphorylated plasmid are ligated together with T4 DNA ligase. E. coli HB101 or other suitable E. coli hosts such as XL-1 Blue

- 20 (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.
- Five µg of a plasmid containing the polynucleotide is co-transfected with 1.0 µg of a commercially available linearized baculovirus DNA ("BaculoGoldTM baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One µg of BaculoGoldTM virus DNA and 5 µg of the plasmid are mixed in a sterile well of a
- Inc., Gaithersburg, MD). Afterwards, 10 μl Lipofectin plus 90 μl Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27° C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27° C for four days.

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After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, supra. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

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To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 μCi of ³⁵S-methionine and 5 μCi ³⁵S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

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Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

25 Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

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Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden),

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pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSport 2.0, and pCMVSport 3.0. Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO)

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Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the marmalian cells are grown in selective medium and the cells with the highest resistance

production of proteins.

Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No. 209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985).) Multiple cloining sites, e.g., with the restriction enzyme cleavage sites BamHI, Xbal and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

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chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the

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are selected. These cell lines contain the amplified gene(s) integrated into

Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

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A polynucleotide of the present invention is amplified according to the protocol 35 outlined in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the vector does not need a second signal peptide. Alternatively, if the

naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

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The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

10 pC6 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene is used for transfection. Five µg of the expression plasmid pC6 is cotransfected with 0.5 µg of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that

confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of metothrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri

20 dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μM, 2 μM, 5 μM, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 -

25 200 μM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

Example 9: Protein Fusions

The polypeptides of the present invention are preferably fused to other proteins.

These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394,827;

Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halflife time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having

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more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.

For example, if pC4 (Accession No.209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

If the naturally occurring signal sequence is used to produce the secreted protein, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

Human IgG Fc region:

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Example 10: Production of an Antibody from a Polypeptide

The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) For example, cells expressing a polypeptide of the present invention is administered to an animal to induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of the secreted protein is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

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In the most preferred method, the antibodies of the present invention are monoclonal antibodies (or protein binding fragments thereof). Such monoclonal antibodies can be prepared using hybridoma technology. (Röhler et al., Nature 256:495 (1975); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J. Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures involve immunizing an animal (preferably a mouse) with polypeptide or, more preferably, with a secreted polypeptide-expressing cell. Such cells may be cultured in any suitable tissue culture medium; however, it is preferable to culture cells in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at 20 about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about 1,000 U/ml of penicillin, and about 100 µg/ml of streptomycin.

The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are selectively maintained in HAT medium, and then cloned by limiting dilution as described by Wands et al. (Gastroenterology 80:225-232 (1981).) The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide.

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Alternatively, additional antibodies capable of binding to the polypeptide can be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody

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whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and can be used to immunize an animal to induce formation of further protein-specific

It will be appreciated that Fab and F(ab')2 and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, secreted protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For in vivo use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art.

(See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP.171496;
 Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

Example 11: Production Of Secreted Protein For High-Throughput Screening Assays

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The following protocol produces a supernatant containing a polypeptide to be tested. This supernatant can then be used in the Screening Assays described in

25 Examples 13-20.

First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) and incubate at RT for 20 minutes. Be sure to distribute the solution over each well (note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The PBS should remain in the well until just prior to plating the cells and plates may be poly-lysine coated in advance for up to two weeks.

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Plate 293T cells (do not carry cells past P+20) at 2 x 10° cells/weil in .5ml DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and L-glutamine (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

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control, one plate of vector DNA lacking an insert should be transfected with each set of minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well. Examples 8 or 9, into an appropriately labeled 96-well round bottom plate. With a vector containing a polynucleotide insert, produced by the methods described in With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. The next day, mix together in a sterile solution basin: 300 ul Lipofectamine

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the even wells, to each row on the 24-well plates. Incubate at 37°C for 6 hours. adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to PBS rinse, and person B, using a12-channel pipetter with tips on every other channel, cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off much time on PBS. First, person A aspirates off the media from four 24-well plates of tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too Preferably, the transfection should be performed by tag-teaming the following

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transfections.

solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock with 1x penstrep, or CHO-5 media (see below) with 2mm glutamine and 1x penstrep. While cells are incubating, prepare appropriate media, either 1%BSA in DMEM

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conical

the incubation period. Person A aspirates off the transfection media, while person B The transfection reaction is terminated, preferably by tag-teaming, at the end of

adds 1.5ml appropriate media to each well. Incubate at 37°C for 45 or 72 hours depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours.

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each well can then be used in the assays described in Examples 13-20 well plate and the remaining supernatant into a 2ml deep well. The supernatants from On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep

ઝ ೫ activity in a particular assay. provides a method of identifying the protein in the supernatant characterized by an proteins, which are then secreted into the supernatant. Thus, the invention further directly (e.g., as a secreted protein) or by the polypeptide inducing expression of other described below using a supernatant, the activity originates from either the polypeptide It is specifically understood that when activity is obtained in any of the assays

HGS-CHO-5 medium formulation:

Inorganic Salts

_	_		_		_	_	_	_			
ZnSO,-7H,O	Na,HPO4	NaH,PO,-H,0	NaHCO,	NaCl	MgSO,	MgCl ₂	KCI	FeSO,-7H,O	Fe(NO ₁),-9H,O	CuSO,-5H,O	CaCl2 (anhyd)
.4320	71.02	62.50	2400.0	6995.50	48.84	28.64	311.80	0.417	0.050	0.00130	116.6 mg/L

Lipids

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Arachidonic Acid	.002 mg/L
Cholesterol	1.022
DL-alpha-	.070
Tocopherol-Acetate	
Linoleic Acid	0.0520
Linolenic Acid	0.010
Myristic Acid	0.010
Oleic Acid	0.010
Palmitric Acid	0.010
Palmitic Acid	0.010
Pluronic F-68	100
Stearic Acid	0.010
Tween 80	2.20

Carbon Source

D-Glucose	Car post pource
4551 mg/L	

Amino Acids

CHINA CHINA	
L- Alanine	130.85 mg/ml
L-Arginine-HCL	147.50
L-Asparagine-H,0	7.50
L-Aspartic Acid	6.65
L-Cystine-2HCL-	29.56
Н,0	
L-Cystine-2HCL	31.29
L-Glutamic Acid	7.35
L-Glutamine	365.0
Glycine	18.75
L-Histidine-HCL-	52.48

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H_2^0	
L-Isoleucine	106.97
L-Leucine	111.45
L-Lysine HCL	163.75
L-Methionine	32.34
L-Phenylalainine	68.48
L-Proline	40.0
L-Serine	26.25
L-Threonine	50.101
L-Tryptophan	19.22
L-Tryrosine-2Na- 2H,0	91.79
L-Valine	99.65

Vitamins

Biotin	0.0035 mg/L
D-Ca Pantothenate	3.24
Choline Chloride	11.78
Folic Acid	4.65
i-Inositol	15.60
Niacinamide	3.02
Pyridoxal HCL	3.00
Pyridoxine HCL	0.031
Riboflavin	0.319
Thiamine HCL	3.17
Thymidine	0.365
Vitamin B ₁ ,	0890

Other Components

HEPES Buffer	25 mM	
Na Hypoxanthine	2.39 mg/L	
Lipoic Acid	0.105	
Sodium Putrescine-2HCL	0.081	
Sodium Pyruvate	55.0	
Sodium Selenite	0.0067	
Ethanolamine	20uM	
Ferric Citrate	0.122	
Methyl-B-Cyclodextrin complexed with	41.70	
Linoleic Acid		
Methyl-B-Cyclodextrin complexed with	33.33	
Oleic Acid		
Methyl-B-Cyclodextrin complexed with	01	
Definal Appropria		

Adjust osmolarity to 327 mOsm

Example 12: Construction of GAS Reporter Construct

One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive responsive element ("ISRE"), located in the promoter of many genes. The binding of a protein to these elements alter the expression of the associated gene.

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GAS and ISRE elements are recognized by a class of transcription factors called Signal Transducers and Activators of Transcription, or "STATs." There are six members of the STATs family. Stat and Stat3 are present in many cell types, as is 10 Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in many cell types though it has been found in T helper class I, cells after treatment with

- many cell types though it has been found in T helper class I, cells after treatment with IL-12. Stat5 was originally called mammary growth factor, but has been found at higher concentrations in other cells including myeloid cells. It can be activated in tissue culture cells by many cytokines.

 The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ('Jaks'')
- The STATs are activated to translocate from the cytoplasm to the nucleus upon tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks") family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarity and are generally catalytically inactive in resting cells.
- The Jaks are activated by a wide range of receptors summarized in the Table below. (Adapted from review by Schidler and Darnell, Ann. Rev. Biochem. 64:621-51 (1995).) A cytokine receptor family, capable of activating Jaks, is divided into two groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and
 - 25 (b) Class 2 includes IFN-a, IFN-g, and IL-10. The Class 1 receptors share a conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a WSXWS motif (a membrane proxial region encoding Trp-Ser-Xxx-Trp-Ser (SEQ ID NO.2))

Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn 30 activate STATs, which then translocate and bind to GAS elements. This entire process is encompassed in the Jaks-STATs signal transduction pathway.

Therefore, activation of the Jaks-STATs pathway, reflected by the binding of the GAS or the ISRE element, can be used to indicate proteins involved in the proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway. (See Table below.) Thus, by using GAS elements linked to reporter molecules, activators of the Jaks-STATs pathway can be identified.

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45		6 6	, 8	25	5	3 15	10	S	
PDGF CSF-1	Receptor Tyrosine Kinases EGF ?	Growth hormone family GH ? PRL ? EPO ? CASSIRF1=IFP>>Ly6)	gp140 family IL-3 (myeloid) (IRF1>IFP>>Ly6) IL-5 (myeloid) GM-CSF (myeloid)	>>Ly6)(IgH) L-7 (lymphocytes) L-9 (lymphocytes) L-13 (lymphocyte) IL-15	g-C family IL-2 (lymphocytes) IL-4 (lymph/myeloid)	II-11(Pleiotrohic) OnM(Pleiotrohic) LIF(Pleiotrohic) CNTF(Pleiotrohic) G-CSF(Pleiotrohic) II-12(Pleiotrohic)	gp130 family IL-6 (Pleiotrohic)	IEN family IFN-a/B IFN-g (IRF)-Lys6>IFP)	ISRE Ligand
.9 .9	nases ?	پر د. د. ^ت		.9 1 1 1	1 1	+ .5 + .5 .5 .5	+ +	+	tyk2
+ +	+	· ‡ '		++++	+ +	. + + + + +	+ ~	• ++	Jaki
+ +	+	+ + +	++ +	2211	, ,	+ -> + + + ->	+ .,	• +·	Jak2
	•		, , ,	+~++	++	+ 12 12 12 12 12	.9 ,	+ 1	Jak3
ພີພູ	1,3	5 5 3 5	00 00	<u> </u>	1,3,5 6		1,3	1,2,3	STAIS
GAS (not IRF1)	GAS (IRF1)	GAS(B-	GAS GAS GAS	GAS GAS GAS	GAS GAS (IRF1 = IFP		GAS	ISRE GAS	GAS(elements) or

6 AAATGATTTCCCCGAAATATCTGCCATCTCAATTAG:3' (SEQ ID NO:3) primer also contains 18bp of sequence complementary to the SV40 early promoter bind STATs upon induction with a range of cytokines (Rothman et al., Immunity 5':GCGCCTCGAGATTTCCCCCGAAATCTAGATTTCCCCCGAAATGATTTCCCCCG sequence and is flanked with an XhoI site. The sequence of the 5' primer is: of the GAS binding site found in the IRF1 promoter and previously demonstrated to Biological Assays described in Examples 13-14, a PCR based strategy is employed to 1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5' generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies To construct a synthetic GAS containing promoter element, which is used in the

with a Hind III site: 5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID The downstream primer is complementary to the SV40 promoter and is flanked

the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is with forward and reverse primers confirms that the insert contains the following digested with Xhol/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing PCR amplification is performed using the SV40 promoter template present in

2

20 ATTTCCCCGAAATATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCC CCCATGGCTGACTAATTTTTTTTTATTTATGCAGAGGCCGAGGCCGCCTCGGC CTAACTCCGCCCATCCCGCCCCTAACTCCGCCCAGTTCCGCCCATTCTCCGC 5': CICGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCGAAATG TGCAAAAAGCTT:3' (SEQ ID NO:5) CTCTGAGCTATTCCAGAAGTAGTGAGGAGGCCTTTTTTTGGAGGCCTAGGCTTT

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30 be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any proteir phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of detectable by an antibody. SEAP, in this or in any of the other Examples. Well known reporter molecules that car With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2

35 element, to create the GAS-SEAP vector. However, this vector does not contain a Xhol, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and The above sequence confirmed synthetic GAS-SV40 promoter element is

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neomycin resistance gene, and therefore, is not preferred for mammalian expression

Thus, in order to generate mammalian stable cell lines expressing the GAS-SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using Salf and Notl, and inserted into a backbone vector containing the neomycin resistance gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning site, to create the GAS-SEAP/Neo vector. Once this vector is transfected into mammalian cells, this vector can then be used as a reporter molecule for GAS binding as described in Examples 13-14.

9

with a different promoter sequence. For example, construction and replacing GAS with a different promoter sequence. For example, construction of reporter molecules containing NFK-B and EGR promoter sequences are described in Examples 15 and 16. However, many other promoters can be substituted using the protocols described in these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, II-2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte.

20 Example 13: High-Throughput Screening Assay for T-cell Activity,

The following protocol is used to assess T-cell activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate T-cells. T-cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12.

Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and Molt-4 cells (ATCC Accession No. CRL-1552) and

Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GAS-SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure described below). The transfected cells are seeded to a density of approximately 20,000 cells per well and transfectants resistant to 1 mg/ml genticin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of interferon gamma. The dose response of a selected clone is demonstrated.

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Specifically, the following protocol will yield sufficient cells for 75 wells containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI

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+ 10% serum with 1%Pen-Strep. Combine 2.5 inls of OPTI-MEM (Life Technologies) with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

During the incubation period, count cell concentration, spin down the required number of cells (10' per transfection), and resuspend in OPTI-MEM to a final concentration of 10' cells/ml. Then add 1ml of 1 x 10' cells in OPTI-MEM to T25 flask and incubate at 37°C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum.

The Jurkat: GAS-SEAP stable reporter lines are maintained in RPMI + 10% serum, I mg/ml Genticin, and 1% Pen-Strep. These cells are treated with supernatants containing a polypeptide as produced by the protocol described in Example 11.

2

On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The exact number of cells required will depend on the number of supernatants being screened. For one 96 well plate, approximately 10 million cells (for 10|plates, 100 million cells) are required.

Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100, 000 cells per well).

12

After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12 channel pipette. In addition, a dose of exogenous interferon gamma (0.1, 1.0, 10 ng) is added to wells H9, H10, and H11 to serve as additional positive controls for the assay.

The 96 well dishes containing Jurkat cells treated with supernatants are placed in an incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples from each well are then transferred to an opaque 96 well plate using a 12 channel pipette. The opaque plates should be covered (using sellophene covers) and stored at 20°C until SEAP assays are performed according to Example 17. The plates containing the remaining treated cells are placed at 4°C and serve as a source of material for repeating the assay on a specific well if desired.

As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate Jurkat T cells. Over 30 fold induction is typically observed in the positive control wells.

Example 14: High-Throughput Screening Assay Identifying Myeloid Activity

The following protocol is used to assess myeloid activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate myeloid cells. Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

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To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 12, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2x10e⁷ U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heatinactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

15 Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl, 375 uM Na₂HPO₄.7H₂O, 1 mM MgCl₂, and 675 uM CaCl₂. Incubate at 37°C for 45 min.

Wash the cells with RPMI 1640 medium containing 10% FBS and then

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resuspend in 10 ml complete medium and incubate at 37°C for 36 hr.

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 ug/ml G418 for couple of passages.

These cells are tested by harvesting 1x10° cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of 5x10° cells/ml. Plate 200 ul cells per well in the 96-well plate (or 1x10° cells/well).

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Add 50 ul of the supernatant prepared by the protocol described in Example 11. Incubate at 37°C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 17.

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Example 15: High-Throughput Screening Assay Identifying Neuronal Activity.

When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1 promoter linked to reporter molecules, activation of cells can be assessed.

Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat phenochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells can be assessed.

The EGR/SEAP reporter construct can be assembled by the following protocol.

The EGR-I promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871 (1991)) can be PCR amplified from human genomic DNA using the following primers:

5' GCGCTCGAGGGATGACAGCGATAGAACCCCCGG -3' (SEQ ID NO:6) 5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO:7)

Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector using restriction enzymes Xhol/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1

25

promoter.

To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

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PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heatinactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

225

Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by growing the cells in 300 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as $5x10^5$

2

Add 200 ul of the cell suspension to each well of 96-well plate (equivalent to Ix10⁵ cells/well). Add 50 ul supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ul of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

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Example 16: High-Throughput Screening Assay for T-cell Activity

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NF-xB (Nuclear Factor xB) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF-xB regulates the expression of genes involved in immune cell activation, control of apoptosis (NF-xB appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

In non-stimulated conditions, NF- KB is retained in the cytoplasm with I-KB (Inhibitor KB). However, upon stimulation, I- KB is phosphorylated and degraded,

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genes. Target genes activated by NF- xB include IL-2, IL-6, GM-CSF, ICAM-1 and

class I MHC.

causing NF- KB to shuttle to the nucleus, thereby activating transcription of target

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Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF-kB promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF-kB would be useful in treating diseases. For example, inhibitors of NF-kB could be used to treat those diseases related to the acute or chronic activation of NF-kB, such as rheumatoid arthritis.

To construct a vector containing the NF-kB promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF-kB

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The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:

S':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO:4)

the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2. (Stratagene) Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

25 3' (SEQ ID NO:10)

Next, replace the SV40 minimal promoter element present in the pSEAP2-promoter plasmid (Clontech) with this NF-xB/SV40 fragment using XhoI and HindIII. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

In order to generate stable mammalian cell lines, the NF-κB/SV40/SEAP cassette is removed from the above NF-κB/SEAP vector using restriction enzymes Sall and Notl, and inserted into a vector containing neomycin resistance. Particularly, the

NF-xB/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with Sall and Notl.

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Once NF-xB/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 13. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 13. As a positive control, exogenous TNF alpha (0.1,1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

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Example 17: Assay for SEAP Activity

As a reporter molecule for the assays described in Examples 13-16, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 µl of 2.5x dilution buffer into Optiplates containing 35 µl of a supernatant. Seal the plates with a plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven heating.

7

Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 µl Assay Buffer and incubate at room

table below). Add 50 μl Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each time and start the second set 10 minutes later.

25 Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

14	13	12	Ξ	ē	# of plates
					plates
80	75	70	65	60	Rxn buffer diluent (ml)
4	3.75	3.5	3.25	υ ·	CSPD (ml)

																													•						
50	49	48	47	46	45	4	43	42	4	40	39	38	37	36	35	34	33	32	31	30	29	28	27	26	25	24	23	22	21	20	19	18	17	16	15
260	255	250	245	240	235	230	225	220	215	210	205	200	195	190	185	180	175	170	165	160	155	150	145	140	135	130	125	120	1115	. 110	105	100	95	90	85
13	12.75	12.5	12.25	12	11.75	11.5	11.25	==	10.75	10.5	10.25	10	9.75	9.5	9.25	9	8.75	8.5	8.25	œ	7.75	7.5	7.25	7	6.75	6.5	6.25	6	5.75	5.5	5.25	5	4.75	4.5	4.25

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Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

potential. These alterations can be measured in an assay to identify supernatants which sodium, pH, membrane potential, or any other small molecule which is detectable by a assay for calcium, this protocol can easily be modified to detect changes in potassium, Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane bind to receptors of a particular cell. Although the following protocol describes an fluorescent probe.

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molecules. Clearly, any fluorescent molecule detecting a small molecule can be used The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small instead of the calcium fluorescent molecule, fluo-3, used here.

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96-well plate with clear bottom. The plate is incubated in a CO₂ incubator for 20 hours. For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash. 12

incubated at 37°C in a CO, incubator for 60 min. The plate is washed four times in the load the cells with fluo-3, 50 ul of 12 ug/ml fluo-3 is added to each well. The plate is A stock solution of 1 mg/ml fluo-3 is made in 10% pluronic acid DMSO. To Biotek washer with HBSS leaving 100 ul of buffer. 2

The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 ul, followed by an aspiration step to 100 ul final volume. re-suspended to 2-5x10⁶ cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml For non-adherent cells, the cells are spun down from culture media. Cells are fluo-3 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. with HBSS, resuspended to 1x10° cells/ml, and dispensed into a microplate, 100

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second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-3. The supernatant is added to the well, and a change in fluorescence is detected. To measure the fluorescence of intracellular calcium, the FLIPR is set for the (6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 8 35

signaling even which has resulted in an increase in the intracellular Ca++

concentration

Example 19: High-Throughput Screening Assay Identifying | Tyrosine

Kinase Activity

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transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase ncluding the PDGF, FGF, EGF, NGF, HGF and Insulin receptor substamilies. In RPTK) group are receptors for a range of mitogenic and metabolic growth factors addition there are a large family of RPTKs for which the corresponding ligand is The Protein Tyrosine Kinases (PTK) represent a diverse group of 2

Activation of RPTK by ligands involves ligand-mediated receptor dimerization, unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and noncytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor esulting in transphosphorylation of the receptor subunits and activation of the e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

12

kinase activity, the identification of novel human secreted proteins capable of activating yrosine kinase signal transduction pathways are of interest. Therefore, the following Because of the wide range of known factors capable of stimulating tyrosine protocol is designed to identify those novel human secreted proteins capable of ectivating the tyrosine kinase signal transduction pathways. 2

with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from 25

0% Matrigel purchased from Becton Dickinson (Bedford,MA), or calf serum, rinsed CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford,MA) are (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or with PBS and stored at 40C. Cell growth on these plates is assayed by seeding 5,000 alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, cells/well in growth medium and indirect quantitation of cell number through use of 8 35

plates can also be used in some proliferation experiments. used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture

Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium To prepare extracts, A431 cells are seeded onto the nylon membranes of

- Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 and a cocktail of protease inhibitors (# 1836170) obtained from Boeheringer Mannheim 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na3VO4, 2 mM Na4P2O7 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example
- 5 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for manifold and immediately placed on ice. To obtain extracts clarified by centrifugation Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum filtered through the 0.45 mm membrane bottoms of each well using house vacuum.
- 5 the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

methods of detecting tyrosine kinase activity are known, one method is described here Generally, the tyrosine kinase activity of a supernatant is evaluated by Test the filtered extracts for levels of tyrosine kinase activity. Although many

- 20 a range of tyrosine kinases and are available from Boehringer Mannheim. PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and biotinylated peptide). Biotinylated peptides that can be used for this purpose include determining its ability to phosphorylate a tyrosine residue on a specific substrate (a
- 25 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl2, 5 mM MnCl2, ATP/50mM MgCl2), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride. order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg2+ (5mM The tyrosine kinase reaction is set up by adding the following components in
- 30 components gently and preincubate the reaction mix at 30°C for 2 min. Initial the reaction by adding 10ul of the control enzyme or the filtered supernatant

The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mm

mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction

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phospotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37°C for one hour. Wash the well as Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of antiallows the streptavadin coated 96 well plate to associate with the biotinylated peptide

tyrosine kinase activity. peroxidase activity is quantitated using an ELISA reader and reflects the level of absorbance of the sample at 405 nm by using ELISA reader. The level of bound incubate at room temperature for at least 5 mins (up to 30 min). Measure the Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and

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Phosphorylation Activity Example 20: High-Throughput Screening Assay Identifying

20 5 substituting these molecules for Erk-1 or Erk-2 in the following assay. Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other kinase activity described in Example 19, an assay which detects activation phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase, phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other used. For example, as described below one particular assay can detect tyrosine (phosphorylation) of major intracellular signal transduction intermediates can also be As a potential alternative and/or compliment to the assay of protein tyrosine

30 25 rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C step can easily be modified by substituting a monoclonal antibody detecting any of the and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp, (RT). The plates are ther Specifically, assay plates are made by coating the wells of a 96-well ELISA

medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants cultured overnight in growth medium. The cells are then starved for 48 hr in basal obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and

႘ filtered directly into the assay plate.

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After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation.

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Example 21: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is be isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer solutions described in Sidransky, D., et al., Science 252:706 (1991).

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PCR products is then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies). The intron-exon borders of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations is then cloned and sequenced to validate the results of the direct sequencing.

PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals is identified by mutations not present in unaffected individuals.

Genomic rearrangements are also observed as a method of determining
alterations in a gene corresponding to a polynucleotide. Genomic clones isolated
according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'triphosphate (Boehringer Manheim), and FISH performed as described in Johnson,
Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is
carried out using a vast excess of human cot-1 DNA for specific hybridization to the
35 corresponding genomic locus.

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Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera

6 (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosomeialterations of the genomic region hybridized by the probe are identified as insertions, deletions, and

10 translocations. These alterations are used as a diagnostic marker for an associated disease.

Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

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A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

Por example, antibody-sandwich ELISAs are used to detect soluble polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10. The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, scrial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbounded polypeptide.

Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbounded conjugate.

Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl 35 phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on

Interpolate the concentration of the polypeptide in the sample using the standard curve the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale).

Example 23: Formulating a Polypeptide

S purposes herein is thus determined by such considerations. administration, and other factors known to practitioners. The "effective amount" for consistent with good medical practice, taking into account the clinical condition of the alone), the site of delivery, the method of administration, the scheduling of individual patient (especially the side effects of treatment with the secreted polypeptide The secreted polypeptide composition will be formulated and dosed in a fashion

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most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If and the interval following treatment for responses to occur appears to vary depending given continuously, the secreted polypeptide is typically administered at a dose rate of to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and polypeptide administered parenterally per dose will be in the range of about 1 µg/kg/day bag solution may also be employed. The length of treatment needed to observe changes continuous subcutaneous infusions, for example, using a mini-pump. An intravenous about 1 µg/kg/hour to about 50 µg/kg/hour, either by 1-4 injections per day or by to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject As a general proposition, the total pharmaceutically effective amount of secreted

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patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers subcutaneous and intraarticular injection and infusion. of administration which include intravenous, intramuscular, intraperitoneal, intrasternal formulation auxiliary of any type. The term "parenteral" as used herein refers to modes to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or intraperitoneally, topically (as by powders, ointments, gels, drops or transderma administered orally, rectally, parenterally, intracistemally, intravaginally Pharmaceutical compositions containing the secreted protein of the invention are

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al., J. Biomed. Mater. Res. 15:167-277 (1981), and R. Langer, Chem. Tech. 12:98-Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (R. Lunger et copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), polymer matrices in the form of shaped articles, e.g., films, or mirocapsules. systems. Suitable examples of sustained-release compositions include semi-permeable The secreted polypeptide is also suitably administered by sustained-release

105 (1982)), ethylene vinyl acetate (R. Langer et al.) or poly-D- (-)-3-hydroxybutyric

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EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA 82:3688-369/ polypeptides. Liposomes containing the secreted polypeptide are prepared by methods (1985); Hwang et al., Proc. Natl. Acad. Sci. USA 77:4030-4034 (1980); EP 52,322: acid (EP 133,988). Sustained-release compositions also include liposomally entrapped

is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal secreted polypeptide therapy. are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes

5 15 known to be deleterious to polypeptides. formulation preferably does not include oxidizing agents and other compounds that are employed and is compatible with other ingredients of the formulation. For example, the carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable formulated generally by mixing it at the desired degree of purity, in a unit dosage For parenteral administration, in one embodiment, the secreted polypeptide is

20 carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl of the recipient. Examples of such carrier vehicles include water, saline, Ringer's uniformly and intimately with liquid carriers or finely divided solid carriers or both. oleate are also useful herein, as well as liposomes. Then, if necessary, the product is shaped into the desired formulation. Preferably the Generally, the formulations are prepared by contacting the polypeptide

30 25 ၓ enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at poloxamers, or PEG. immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as the dosages and concentrations employed, and include buffers such as phosphate, polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates. manose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, The carrier suitably contains minor amounts of additives such as substances that

concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of The secreted polypeptide is typically formulated in such vehicles at a

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about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

into a container having a sterile access port, for example, an intravenous solution bag or Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are placed Any polypeptide to be used for therapeutic administration can be sterile. vial having a stopper pierceable by a hypodermic injection needle.

formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution, and the Polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized lyophilized polypeptide using bacteriostatic Water-for-Injection.

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pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of present invention may be employed in conjunction with other therapeutic compounds. The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical

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Example 24: Method of Treating Decreased Levels of the Polypeptide

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administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an pharmaceutical composition comprising an amount of the polypeptide to increase the It will be appreciated that conditions caused by a decrease in the standard or increased level of the polypeptide comprising administering to such an individual a normal expression level of a secreted protein in an individual can be treated by activity level of the polypeptide in such an individual.

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polypeptide is in the secreted form. The exact details of the dosing scheme, based on For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the administration and formulation, are provided in Example 23.

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Example 25: Method of Treating Increased Levels of the Polypeptide

a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer. present invention. This technology is one example of a method of decreasing levels of Antisense technology is used to inhibit production of a polypeptide of the

2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, For example, a patient diagnosed with abnormally increased levels of a provided in Example 23.

Example 26: Method of Treatment Using Gene Therapy

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hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to separated into small pieces. Small chunks of the tissue are placed on a wet surface of a expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and issue culture flask, approximately ten pieces are placed in each flask. The flask is urned upside down, closed tight and left at room temperature over night. After 24 One method of gene therapy transplants fibroblasts, which are capable of he bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS,

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penicillin and streptomycin, is added. The flasks are then incubated at 37°C for approximately one week. 2

At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The nonolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is erminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and fractionated on agarose gel and purified, using glass beads. 25

using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear The cDNA encoding a polypeptide of the present invention can be amplified backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to 8 33

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transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is being produced.

The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

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It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference.

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\$ 6 35 છ 25 20 5 5 S (iii) NUMBER OF SEQUENCES: 644 (ii) TITLE OF INVENTION: 186 Human Secreted Proteins (i) APPLICANT: (v) COMPUTER READABLE FORM: (iv) CORRESPONDENCE ADDRESS: (vi) CURRENT APPLICATION DATA: (E) COUNTRY: USA (C) CITY: Rockville (D) SOFTWARE: ASCII Text (B) COMPUTER: HP Vectra 486/33 (A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage g (B) STREET: 9410 Key West Avenue (A) ADDRESSEE: Human Genome Sciences, Inc. (B) FILING DATE: March 6, 1998 (A) APPLICATION NUMBER (C) OPERATING SYSTEM: MSDOS version 6.2 (F) ZIP: 20850 (C) CLASSIFICATION: STATE: Maryland Human Genome Sciences, Inc. et al

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(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE:

(1) MARE A A Addra Brookeride (1) Brookeride (1) Brookeride (2) Brookeride (3) Brookeride (4) Brookeride (4) Brookeride (5) Brookeride (4) Brookeride (5) Brookeride (6) Brookeride (7) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (8) Brookeride (9) Brookeride (10) Br						09	98				72			09
(viii) ATTORNET/AGENT INFORMATION: (1a) RESISTANTION NUBERS: 56,373 (c) REFERENCE/DOCKET NUBERS: 56,373 (c) REFERENCE/DOCKET NUBERS: 56,373 (d) TELEPHONE: (301) 309-8504 (e) TELEPHONE: (301) 309-8504 (e) TELEPHONE: (301) 309-8504 (f) TELEPHONE: (301) 309-8504 (g) TELEPHONE: (301) 309-8504 (h) TELEPHONE: (301) 309-8504 (ii) SEQUENCE COMMENTARY RECOGNING CONCOURTER (iii) SEQUENCE DESCRIPTION: SEQ ID Not: 1: GOGRANCOA OCCUMANTY TOTACADAM CONCOURTER (iv) TOTACADAM CONCOURTER ADMONITORY CONCOURTER (iv) SEQUENCE RESISTANTION: SEQ ID Not: 1: GOGRANCOA CONCOURTER TOTACADAM ANCOURTER ADMONITORY FOLOCOTICA CONTINUENT ADMONITORY CONCOURTER 100 TOTACADAM ANTONIORY CONCOURTER ADMONITORY ANTONIORY CONCOURTER ADMONITORY CONCOURTER 100 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 TOTACADAM ANTONIORY CONCOURTER 101 T	 (2) INFORMATION FOR SEQ ID NO: 2:	iĝi Si	E DESCRIPTION: SEQ ID NO:		SEC	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 3: GOGCCTCGAG ATTTCCCCGA AATCTAGATT TCCCCGAAAT GATTTCCCCG AAATGATTTC	CCCGAAATAT CTGCCATCTC AATTAG	(2) TANDBHANTON POB SED ID NO. 4.	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 27 base pairs		FION: SEQ ID NO:	(2) INFORMATION FOR SEQ ID NO: 5:	SEQUENC (A) (B) (C) (C)	(xi) SEQUENCE DESCRIPTION: SEQ 1D NO: 5: CTCGAGATTT CCCCGAAATC TAGATTTCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG
(viii) ATTORNEY/AGENT INFORMATION: (a) NAME: A. Anders Brookes, Esq. (b) REGISTRATION NUMBER: 36,373 (c) REPERENCE/DOCKET NUMBER: PS002.PCT (vi) TELECOMMUNICATION INFORMATION: (a) TELECOMMUNICATION INFORMATION: (b) TELEFAX: (301) 309-8439 (c) INFORMATION FOR SEQ ID NO: 1: (d) SEQUENCE CHARACTERISTICS: (a) LEARTH: 313 base pairs (e) TRES: INCHESTESS: dauble (ii) SEQUENCE DESCRIPTION: SEQ ID NO: 1: GOODATICOROL GOODANCT TOTACADAM COLOURIDE COACHORITE ANTICALAGG ACCOLANTE TOTACADAM COLOURIDE COACHORITE ANTICALAGG ACCOLANTE TOTACADAM COLOURIDE COACHORITE ANTICALAGG ACCOLANTE TOTACADAM ACCOLANGA ACCOLANTA TOTACAGGA ACCOLANG ACTICATOR TOTACADAM ACCOLANGA TOTACAGGA TOTACATOR GACGOCATO ACCOLANGA ACCOLANGA TOTACAMACA TOTACATOR GACGOCATO ACCOLANGA ACCOLANGA ACAMAACAN CHORACACA ANGEGIAG TOTACATOR ACCOLANGA ACAMAACAN CHORACACA ANGEGIAG COCAMAGACA ACCOCANTOR ACAMAACAN CHOCANACC ANAGACAG TOTACATOR CHARGATER ATCOLAGGA GAGACAGA ACAGACAGA COTACATOR COTACATOR ACAMAACAN CHOCAGA ACAGACAGA TOTACACATOR COTACATOR ACAMAACAN CHOCAGAGA COCAMACATOR TOTACATOR COTACATOR ACAMAGAGA CHOCACAGA GAGACAGA COTACATOR COTACATOR ACAMAACAN COCACAGA GAGACAGATOR TOTACATOR COTACATOR ACAMAACAN COCACAGA GAGACAGATOR TOTACATOR COTACATOR ACAMAACAN COCACAGA GAGACAGATOR TOTACATOR COTACATOR ACAMAACAN COCACAGA GAGACAGATOR TOTACATOR COTACATOR ACAMACACACATA CACACAGAGA ACACACAGA COTACATOR ACAMACACACATA CACACAGAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAGA ACACACAC		8	<u>0</u>	15	20	25	30		35	. 40	. 45		50 50	09
	(viii) ATTORNEY/AGENT INFORMATION:	(A) NAME: A. Anders Brookes, Esq. 5 (B) REGISTRATION NUMBER: 36,373	(c) REFERENCE/DOCKET NUMBER: PS002.PCT 10 (vi) TELECOMMUNICATION INFORMATION:	(A) TELEPHONE: (301) 309-8504 15 (B) TELEFAX: (301) 309-8439	20 (2) information for SEQ ID NO: 1:	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 713 base pairs (B) TYPE: nucleic acid (C) STRANDELNESS: double		•		TCAAGTTCAA CTGGTAGGTG GAGGGGGTGG AGGTGCATAA TGGCAAGACA AAGGGGGGGAAGAAGAAGAAGAAGAAGAAGAAGAAGAA				

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	AAGCCAACTG AGATACCGTG ATGGTGTTGA TTTCTTTCAA TGATGCTTAC CATCTATTT	480	ACCC	ACCCINGCAM ITTOGGOCCTG ACCOGGTOGA TCACGAGGTC AGGAGATCGA GACCATCCTG	420
ų	AGCCACTGAG CCTTTTATTA TYTOTCTATT TGTAAAGTTT ATTTGTCTTA ACTCATTTAA	540	GSTA	GSTANCATIGG TGANACCCCG TCTCTACTAN ANATACANAN ANANANANA ANAN	474
•	taaathtact gtttatctgt ttctgaaaa aaaaaaaaa aa	582			
9			(2)	(2) INFORMATION FOR SEQ ID NO: 14:	
2	(2) INPORMATION FOR SEQ ID NO: 12:			(i) SEQUENCE CHARACTERISTICS:	
	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 665 base pairs		:		
15	(B) TYPE: nucleic acid (C) STRANDECNESS: double (D) TOPOLOGY: linear		15	(D) TOPQLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:	
6	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:		TTATI	TIATOTIOSO GAGCAAGACC TGATAGCCAG CCTTTACATO GGAGTATAAT TCTGTCCTCC	09
2	OTTROGOGOT GACCOCCACC TGCTCCCCCC CCCCAGGACA CACCTACTCC	09	-	ATCTCATAAG CCCCAGTACC TGAGCCAGAA TGATTATAAC CAACCACACT GTCTCTTTAT	120
	CACCACCACC GCCCTGGCT ATGATGTTCC TCACCCAGGG CGGCCTCTG CCCTCTACTC	120	CATO	CARGGATGGC TITNAGCAGTA GGTTATTITC ARCATTGCCA ITTGTAGCTC TACAGGGGTT	180
25	STOCCAGGCC CACTTOCCAG GCAGGAGCC TCCCCAAGCC TTCAGGGCTG CTCGGAAGTCA	180	25 TATA	TATAGTAATT TETECHETT TAAGTETETE CETEAGTGCE TOTTOTTATE AAACTEATTG	240
	CCTOTTIGGIA TGGACTIANA GGACCCTTGT GTGGGAACAG GTGCTCCCCA AACACCCTGC	340	CIC	CTCTCTCANG CAGTTCAGCT CTCCATTCTC CCTTATCGGG CAGACCTCTG TTGCAGAGAG	300
ç	TISCHIBACTIBC CHABACC CTCTIBANDA GNABODOCNA GNCTCCCTO	300	30 AGAA	AGNATATIVAC TTCC	314
ટ્ર	GACCCCTGCA GGGCAGGCAG CTTGGGCCCG AGCCCAAGCA TTTGGCTCTG CTGCCCCCAA	360	S		
	GOGGACAGGA AGCOTOTIVOS GCOTOTITOCO TITOCITOGACA AGGOCOCOTIS COTITIGOCITO	420	5	AND THE TOTAL TOTA	
35	ACATAAACTG TACAGTATTT TCATTAAAAG CCTCTTTCAT AAAAA	465	35 (%)	(a) INFORMATION FOR SOLUTION (a)	
9 9	(2) INFORMATION FOR SEQ ID NO: 13:		40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 613 base pairs (B) TYPE: nucleic acid (C) STRANDENESS: double (D) TOPOLOGY: linear	
	(1) SEQUENCE CHARACTERISTICS:			(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:	
Ý	LENGTH: 474 by TYPE: nucleic		d5 crcs	CTCATATTOC COTCTOCCTA AAAGTGAACA TOCCATTGAT CAATCTGCTT TTATTATATT	
5	(C) STRANDEANESS: GOUDLE (D) TOPOLOGY: linear			ATGTTCCTAA TOOTCOCCAAG CAAGACAAGA AOTAGAAAGA AAGATGOTOT AAGCTCAAGA	120
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:		ACCC	ACCACTAAA TCTATCCTAT GGCCTGGGTT CACCCAGCCT GCTTTGTGGGA TTTTGTCTCA	180
20	ATISCIAATTICC TISCICAÇÃOS CITTICITOTIOS GIBCCIACITO TISOCICITITIS TISATISFICACO	09	50 cmar	CTATAACAGA GCTCCCAAGG AGACTGCAGA GTCAGCTCCC TTAAGCACTG TAACTAAAGC	240
	ATATCCCTAG GCTTCTCCCC CTCCTAGAAG GCCTTCTTGA TAGATTAGAA AATAAGAATG	120	ST.	CTAACTICTIC COTTCCACCC AACAATGTYC CCAGCTCATC CTCTTTCCCR AAGTCCCCTT	300
y	AGTGACATTT CCTATGTGCA TATAAGAAGG AGCCACAAGA CATGTCTTTT AAATAAAAGG	180	TCTG	TCTGCCCCAG ATGCGAATTG CATTTAACTA ATCCTCAAGT GAAATGTCCA CACAGRATTC	360
2	ACAGIOTICA TECTITITAGE TOCCOANTAG AACCITOGIC TEATECTECT GGAGCIAGGE	240		CAPTITIANT AGCATACCAT AGTITITIGIG CAANITIGCT TICAGARGAC TCCCAFTGCA	420
	CITIAAAACA GCITCIGIGI ITCTCAITIG TCTCAGIGIT ITGCCAGGGI ITTAITCGGAA	300	SCTO	GCTGCTCAGA GACGCTAAMG GCAGGCCTC TTGAMGCTTT CCCGATAGCT TTCAGCTGCA	480
9	AGATAATGIT COSTITAAAA TAITITCCIAA TGAGGCGGGG COTGOTGGCT CACGCCIGIA	360	60 ATAG	ATACCTETTA GGCAGAATGC CATGAGGGTC CTGCCCAACT GTATTACTGG GGAACACCTG	240

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(2) INFORMATION FOR SEQ ID NO: 18:

	TACTGAACTA TCTTTATGAC TTTGGATTTG ATCAGAGGTT TAAAAAAAAA AAAA	CACTATOSTE TECCTECTOT TEACAATOST ATTIACAGGA GACCTIGICA TEAGAGGACG	TCTCCCAGAA AAAAGTACTA CAACAGGCTC AAGGGATATG CTTTGGTGGT CAAGGGATTA	TITIOCATATIC CACTOCAGIT CIGICACCAA AGATTITIAAT CITICAGANGG CAATTITOCIC	ATTCACATTA ACTOTOCTAG GATACTTCTC TIGAGGCITI GGAAAACTTC TICCTIGAAA	GNICIACATTO ACTOCIACCIA ACCIAAGCTA TAGAAAGAAA TGATTGACTT TTTAALATAT	GAAACTANAT CCCGGGGCTT TIDACNOGTA CTTGGGAAAT AAGTATTGGG TAATCACTAA	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 17:	(C) STRANDEDMESS: double (D) TOPOLOGY: linear		(1) SEQUENCE CHARACTERISTICS:	(2) INFORMATION FOR SEQ ID NO: 17:	СССТСАНАНАС СПАЛІТІСТС СПОТІТАССЬН АБВИССИСНО ОООВИЛАННЯ АНАННЯ	GGGAAGCAT TIGGICCIGG TINIGITINI TACAACAICA TIGCACICIG GGACICCAGI	GTEAGTGGTC TOCCTGCCCA AGGAGCCTGA TTGGTGGGAA ATGGCATCAT CTAATATGAT	TENTETTAG TECHARIGA CCAGGETETCC CCCCCACAGC CECTGETCIGG TCCCTCATIG	осствотого позитесска спестотное маменовско смоссамисм поммососсе	OCCOCCOAT TOMACCOTOG GOTOTOMAMS TITITIGCOTO TOTOGOTCOT TOTOTOTOGO	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 16:	(C) STRANDELMESS: GOUDLE (D) TOPOLOGY: linear			(2) INFORMATION FOR SEQ ID NO: 16:		CCNANTATO CAC
	414	360	300	240	180	120	60						356	300	240	180	120	60						;	613
	?	55		50		į	45		40		Ę	a n	30		25		ţ	20		15		Š	10	_S	
алаалалаал	ATGAGCCC	TOTOGA	AGGAG	ACTG	8	>																			
	ATGAGCCGAT GGGCCCTGGA GGCAGCCCAT TAAAGCATCT GGCTCGTTTT TGGAAAAAAA	TOTOGRAFIOC TORROCICARC CRARGOCIOG CARGOTOTOG OCCICATITA ROSGRITICIO	навнесттве тнегессени осснанаесе итенасеная сетветотан осснествее	ACTOGROUAG GRAGOTIGAG TOGGAGCTICA GROCTAGAAG GROCTIGAAG ARCTIGGACTIG	GGOCKTTATC CCAGGAAACT TRATGITTIC TAGAAGCTAA GCAGCTGCTG GGACTCAGGG	AGAGECCEAG CENTRETTA ATTAKANCA GIGETYCCET GAACTGCETE CCCCACCCET	отвежноськи вежноствам асторовтое ттотпетвор сетветвето есвесансее	тастосьное тованостоее тованованое тованстоеет остоснале оссостането	CCCCCCCCC CCCCACACT TICHGGAGTC ACCCCCCAGC ATTIGGGGTT GGGTTGGCCC	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 19:	(D) TOPOLOGY: linear		(2) INPORMATION FOR SEQ ID NO: 19:		ACTICICIST ATTETTOCIC ATACTTOCCT ACCCCAAAT TAATATCAG	GATOTICOTOT TOGAGOGATO AAGATICAAGT TATICCTCTIC AGAAATTCCT AGACCCCTTC	GASTINCCTS AMOSICCAAA CCICAGAACC CICOOGCACC TGAGAGAGAS GTITTGCGGC	оссманалос аталагалос оссманасет тогомаллал останасеста аламалалсяс	GTTGGGCCTG CTTGCCGCGG GAGTGACGCT ACTTCTTCCA GAGACCAAGG GGGTCGCTTT	CITICARAGIC TICAGGCIGA GGGAGGICIG GCAAGCCITG CCCCICATIT IGITIIGCGGI	CAGGAACYTC GGAGTGATGG TGTGTTCCTC CCTGTGTGAC ATAGGTGGGA TAATCACCCC	AATCACCATT GCAATACAAA TGATCTGCCT GGTGAATGYT GAGCTGTACC CCACATTCGT	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 18:	(B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 469 base pairs

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		ACILMASTIC AMAMAMATAT ACATAAATAA GATAAAGCIG ACCIGTAGAT ATAGCAGGTT	395
(2) information for SEQ Id No: 20:		ATAAAGCTTA GAGTIGICTA AGTIGAGIGC AAATTITICCT CIGAICTTITC TGAIGCGGAA	420
(1) SEQUENCE CHARACTERISTICS:	2		
(A) LENGTH: 741 base pairs		CANANAAGCA GICATGITIG TIATGICATT GGAATGGAAC CCGAGAAGAG AGCATGCTGT	480
(B) TVPE: nucleic acid		STICTIONS GACAGGAAAG CINSCONSCA CCAAGNCHGA ACCACCA TCANGGRAA	540
(b) Topology: linear			
-	01	ATAGATTATG TGCTGGAACA TAITTTCACAC CGGCCTGGCA GTAAACACTT GTAGTGTTGT	909
(X1) SEQUENCE DESCRIPTION: SEQ ID NO: 20:		GCAGTGGAAA CGSTCATCTT CCGCTAAAGC ACGCCGTGTT GTGCAGCGGA AATGGTCATC	099
tcttgaagag tgtacagtac aggattatta taatgaaagt ttatatcaac aggstttcgf	09	TOCTOCTAAA ACACAGCTTC CATOGTAATG TATGCTCCTT ACTCAAAGAG TGTGGTCCCA	720
Toscictisca tatattataa gcaaaagaa ttostaaast gccacastat tccasataac	120	AACAGCCTTT GGGAGGTCCT CCTTGATTCA TGGATGAAAC CTGGAACATC TTGAGGACTG	780
TITICAGITG COSCULTOR POTCOTICIT TAAITIGAAA CCIAGATACA TGCAGTAAAA	180	ASTINACCAT AGGICCTINA ATAACTCTCC ACAGGITITIT CITAGITITAS CICTACANG	840
ACTAGGAGAA TGACTTTTAG CCTTGGGGAC AGCCAAGTTT TGTTGATAAA CCTATTTCCT	240 20	AGGGTGTGCA GCAGCCTGTT CAAAGTCATA TTTTCTGGGA AATATTTCCA GTGTTTATTT	900
AGCATGCCTT CAGGAAGTTG TGCCAGACCC TAGATTGTGA AGGACCCACT GTTCTTCTGT	300	GCACTITIAGC CCACTCTGTG TAGCCTTAIT TCTTCTAAAC TCACCATTAA TCTGAATAAT	960
TOTACGAGCT COCTGAACCA TIGITCAGAG GACCAATGTC ACATCGCTTC ATGGGCATGG	360	ACTICAAATTT AGGGGGACTG TATTTGCCTT A	196
NCCATOGGAG CATCTGGGTG ATAYCTGTCT ACAGTATTGG CTCTTCTGGG AGGCTGATAC	420 25		
ACAAGGCTC TCTTCCACAT GATCATTTGC AAACCTCCCC CAGCCCCTAC CATCCAATGT	480		
GGNAGGNANA CHAGAACTOC CTGNAGAGA GTOCDAGCTA CAGNINGACA GCOTGTGCAT	30	(2) INFORMATION FOR SEQ ID NO: 22:	
TOCGOCTOTC ACCTTCCTC TOCCACTTCT GTATOCTCAG AGATGCTGCG TGGATGTTC	009	(1) SEQUENCE CHARACTERISTICS: (A) LEXYTH: 651 base pairs	
CITAACCICA GCTGACITCC CIGIGAATGT CTAATGCTAG ITCAGGGCCT CCAGGCAITG	. 099		
ATTIGIACAG TGGTAACTCC CAATGAGGCT TCTGTTATCA TTTGGTGTGC TTTYTCTGTC	720 35	(D) TOPOLOGY: linear	
ATTABABGAA ATGATTTTCC C	741	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:	
	04	CCACGCOTCC GGAATTCCCC TGAGGATCTT GGGCTATCTT TGACAGGGGA TTCTTGCAAG	9
	?	TTGATGCTTT CTACAAGTGA ATATAGTCAG TCCCCAAAGA TGGAGAGCTT GAGTTCTCAC	120
(2) information for SEQ id no: 21:		AGAATIGATO AAANGGAGA AAACACACAG AITGAGGATA CGGAACCCAT GTCTCCAGTT	180
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 991 base pairs	45	CICARITICIA AMITIGITICE TICCIGAAAMI GATAGIANICE TGATGAANICE AGCACAGGAN	240
(B) TYPE: nucleic acid			
(C) STRANDEDKESS: double		GSIGAASIAC AACIGASICA GAATGATGAC AAACAAAGG GAGATGATAC AGACACCAGG	300
TOLOGOT: TYURBUT	S	GATGACATTA GTATTTTAGC CACTGGTTGC AAGGGCAGAG AAGAAACGGT AGCAGAAGAA	360
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:		OTITICIATIVE ANCICACITIG TGATTCGGGG AGTCAGGCAG TTCCGTCACC AGCTACTCGA	420
GGCAGGAGTC TCCCCTGGGG AAGTTTTCT TTTTCAGGAG GGAGAGGGC TTTCCCAGGT	09	TCTGAGGCAC TITCTAGTGT GTTAGATCAG GAGGAAGCTA TGGAAATTAA AGAACACCAT	480
AATOTOTOTA GAGTOTTGGG CAGAAAATOT GGGACCACAC CACACCAGTT CTCTCCTTAA	120 55	CCAGAGGAGG GOTCTICAGG GTCTGAGGTG GAAGAAATCC CTGAGACACC TTGTGAAAGT	540
TCCAGGICAT TIGCCITCTA TCCCAGGIAT GITTCCAGIG TCCICTGGGT GITTCCAAGA	180	CAAGGAGAGG AACTCAAAGA AGAAAATATG GAGAGTGTTC CGTTGCACCT TTCTCTGACT	909
GCAACAAGAA ATGAATAAAT CTCTGGTGAG TTGTTTATTT GTTCTTCACT TTGTTTTACA	240	GAAACTCAGT CCCAAGGGTT GTGTCTTCGG AGGCATCCAA AAAAAAAAA AAA	653
CICTAITITIC TCACHTAIG GOICICIOIG AATIAAAAG GAAAAGIAGA AATAAGIAAA	09 000		;

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TOGATOCCCT GACTGAGCTG CITAGCACTG CCCTGGGGCC GAGGCAGCTT CCAGAACCCA TGGATATCCA GTATACCAAA GAGATGAAGG GCTGGGCACT GGTGCTGGCA SGARCTGGCA TTGGACTCAT GGTGCTGCAT GCAGAGATGC GCTGGCTAAA GTGGGKAGGC CTTGGCCCAC CTGAGGCCCC AGGTGGGAAC ATGGTCACCC GOTOGTOGAC ATCOTCTCTG CCACTCCTGA CCCAGCCCTG AACAAAGCAC CTCAAGTGCA GCCAGCAGTC CAAGTAGCTG GACCCACGAG GAGGAACCAG GCTACTTTCC CCAGTACTGA CCGTGGTGGC CCGGAAGCTG GAGTTTAACA AGGCAGAGAA GCACGTGCAC AACTTCATGA GCAAGATCGT YTGCCTGTGC ACTGGAGTCA TGGGTGTCTG CTGCACAGCC CTGCTGGTGG TITOSCCITOGO GOCCITICAGA COCCGAAAGC OCTITOCITOGA OCAGGAGAAG ICTCIROCCO CCTRRRAGA AAGCTRRACTG AGCCCCAAGA CCCCRRRAGGC CATRRRAGCRR GATCTRRFTRC GOCAGOCTIGA CGACCTOCAA GCCACAGTOG CTGCCCTGTG CGTGCTGCGA GGTGGGGGAC AGGACCAAAG GOGGCCCTOG CTTGGAGTGG GTTGGCTTGC TGATGGCTGC TGGAGGGGAC AGAATOTGAG CAGCTCACAC COGGCCCTGG AGAAACAGAT TGACACGCTG GCGGGGAAGC AACAAGTGAA CTCCATGGTG GACATCTCCA AGATGCACAT GATCCTGTAT GACCTGCAGG ANCTOCTOGO COCCATOAAC OCOTTOCOGO AGGTGCOGOT GAAACACOGO AAGGTCOGG GGATOTICTA CAAACATACT CGCAGGAAGG AGTCTCATGC TGCCCGCANG CATCAGCGCA TGATICCCCAT CACATTCCTG ACCATCGGCT ATGGTGACGT GGTGCCGGGC ACCATGTGGG TOTOGTTCOG GOGGTOCTCG OCTOTCANTO CCACTGGGCA CCTTTCAGAC ACACTTTGGC INFORMATION FOR SEQ ID NO: 23: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: TACCCTCATC AAAAACACTC TCACTATGCT GCTATGGACG GATCCAGGAT TCTGGGAGGC TTCAGTTACC GCTGGCCGAG CTGAAGAACT AGTICCAGGCG ACTIGGAGGCA GGACTCCTIGG GTCCCTIGGGA AAGAGGGTTAC SEQUENCE CHARACTERISTICS: AGAGCTOCAG AGAGCACCTG GTGGGGAGGA AGAAGTGTAA CTCACCAGCC CTGGGGCGGG GCTGGAGGTG GCGCCCCCTG GTGGGACAAC AAAGAGGACA (A) LENGTH: 1486 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AGTOCGCTGC CCGAGTGCTA CAAGAAGCCT 23: ACCTCCAGCT 1140 1380 1320 1260 1200 1080 1020 960 900 840 780 720 660 600 540 480 420 360 300 180 120 8 20 8 SS 50 45 ઝ 30 25 5 5 S TATATTTTCC ACCARGICTT GACAACCIGT ACTCTTCCAA TOTCATTTAT CAGITGTAAA TACCITITICT ATTITITATION TENGATECET TOTATENANT TECTTOGRAM TONISCRITT CAGAATCCCC CAGATGGCTT CATGGCCCCC AAAGCATGGA AGGTCCTGAC AGATTATTAC TIGICICCA TOCCATOCCI GITAATICCI GGATGGGGAC AAIGICCOTC AIGGCCITAA GICTAATAAG AGAAGICTIA AIGGCCICIG IGAATAAIGI AACICCAGIT CTARARGAGG TOTOGCTCAC ATCARGATTC TTCCTGATAT TTTACCTCAT GCTGTACAAA GOICCCAGAT TICTIAAGGC TITGTITICAC CATGTGTCTA GITACITGCT GAAAAGTGAA TICTIATACA TITCATAATA AAATTAGCTC TAIGTATITT CTACIGCACC TGAGCAGGCA ACACACAT ATACATACAA AGTCAAACTG AAGACCAAAT CTTAGCAGGT AAAAGCAATA TRATCITITA TRATCAAGTA AAAGTTOTOT CIRTAATIRA AAAAATRIRI ATRIATATAC AGGICCCIGG AGAAGAACTA AGCCITIGGT CCAGAGITTIC ITTICIGAAGI GCICITIGAT GUAGGAATAA GAGAGGGCAA GTOGTTOGAA CAAGGGGTGG GTTCCGAGGA TGTACCGTGT CONTRAGON CICCOICOG ANCGANGIGO TOCCOCOGCO GCCGCCGCCG TCCCGCGICC CTTCGCCGTT TCTCCTGCCA GGGGAGGTCC CGGCTTCCCG TGGAGGCTCC GGACCAAGCC 3 AACTOGAGGG GGGCCCRKAC CCAATCWCCC TATAGTAKAC GTANNN GCCTTAATGT TGTAATCATA TCTTACGTGT TGAAGACCTG ACTGGAGAAA CAAAATGTGC ATATATCAGA TOTOTCCTCT TCTGTACAAT TGACAAAAA AAAAATTTTT OCCIDANCAGO ICICICICOS CCIOCGAMAMA ACAMOGATAM CITITICCCCI GGAGGAGIAC TITTAATAGC ATACAGTGAT TIGATGAAAG GACGICAAAC AATGIGGCGA TGICGIGGAA AGCAAAAGAC GGAGAACCAG CAGAAATCCA CCAATGTAGT CTATCAGGCC CACCATGTGA TICGGICICI OCICCOGGGA CCCGGCICOG CGCAGCCAGC CAGCATGICG GGGAICAAGA AGITATOTT COOCTOTT GOTGTGGTCA TIGTGTCTTG CAGAAAGGAT GGCCCTGATG AATAACOIGA ATTITATCIT AGAGATCIGT INFORMATION FOR SEQ ID NO: Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLOGY: linear (A) LENGTH: 2323 base pairs 24: GCAGCCTATT TCTGTCACAA

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ACACGGTGAC AAGTTATATT

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	CHECHOCHEC GOCHECTETA ATRABABATA ATTCACACTA TCAGACTASC AMBOCACTAS	. 1380	
	AACTIGGAAAA GACCACAGAA AACAAAGAAT CCAACCCTTT CATCTTACAG GTGAACAAAC	1440	
2	TOTGATGATG CACATGTATG TOTTTTGTAA GCTGTGAGCA CCGTAACAAA ATGTAAATTT	1500	S
	OCCATTATTA GGAAGTOCTG GTOGCAGTGA AGAAGCACCC AGGCCACTTG ACTCCCAGTC	1560	
9	TIGOTOCICATO TICTACACOGO ACCAGOGOTOA GATTOCOCTIC ACCTECTIVA	. 1620	9
2	CAAAGITOCT CGAACAGAAA GTOCTTACAA AGCTGCCTTC TOGGATACTG AAAGGTOGAG	1680	:
	TITICIGAC TOCACTGATT TTATTGCAGT TGAAAAAAA AAAAGCTAT TCCAAAGAIT	1740	
15	TCAACCTOTT CTGAGACATC TTCTGATGGC TTTACTTCCT GAGAGGCAAT GTTTTTACTT	1800	13
	TATGCATAAT TCATTGTTGC CAAGGAATAA AGTGAAGAAA CAGCACCTTT TAATATAG	1860	
Ś	OTCTCTCTGG AAGAGACCTA AATTAGAAAG AGAAAACTGT GACAATTTTC ATAITCTCAT	1920	70
3	TUTTAAAAAA CACTAATUTT AACTAACAAA AGTTUTTTTG AGAATAAGIT ACACACAATG	.1980	1
	GCCACAGCAG TITGCCTTTA ATAGTATAGT GCCTATACTC ATGTAATCGG TTACTCACTA	2040	
25	CTCCTTTAA AAAAAAAAC CACCATATT ATTGAAAACA TGAGACAGA TTATAGTGCC	2100	22
	THANCCGAIA INITITICIGA CITAAAAAT ACAITIYAAA CIGCICITCI GCICIAGIAC	2160	
S	CATGCTTAGT GCAAATGATT AITTTCTATGT ACAACTGATG CTTGTTCTTA TTTTAATAAA	2220	ಜ
ર	tttatcagag tgaaaaaaa aaaaaaaaa aaaaaaaa aaaaaaaa	2280	
	AAAAAAAAAA AAAAAAAAAA AAAAAAAAA AAAAAAA	2323	
35			35
	(2) Information for SEQ ID NO: 25:		
40			40
	(A) LENGIN: 003 DASP PALES (B) TYPP: nucleic acid (C) STRANDELNESS: double		
45	(D) TOPOLOGY: Linear		45
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 25:		
	GGCACGAGCC TOTGTGGTCA TGTTCCTCGT GGTGCAGTAC CTGACATGAG CCAGCCACGC	09	
20	TCAGTGOCTG AACAGCATTC CCACAGCCTG CAAGTGTGTG TGTGTGTGAA AGAGAGGG	120	8
	GGGCCCAGAG CCGCCTTTTG AAATGTTTGC CTGTCTGAAC TGTGAAGACA CTTGGGAATG	180	
55	ATTGTGGTCT AATTTCCAAC CTGCTCTGTT TTCTGTGACA TCTTGGAGGG GAGCTAGTGC	240	55
}	CACACCATGG GOGGIGGITA GAAATGAAAA AGTCCCGGGT CTGTCTCTCT CACTCTCGCT	300	
•	CTCATGGGGG AGGGAAGAA TGGCTFTGGT GGCTFTGTTC ACACAGCTGA TGCGTGCTGG	360	
9	GAAGGIGICC ACAGIGAGCC TGTGTGCAGG ACTGTGCACA CGGTTCACAC TTGTCACAT	450	8

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480 540 9 099 683 120 180 420 540 9 99 720 780 840 480 CTCACAGCCA TTATATTAAA TAGTAGGTCG ATTCACATCT CGTGCTCCTG GCCACCTTCC CCTGTGCCTC AGTGACATGT AGATGACTGA CTGCCAATAC TTGTCACCAT TCCCTGGAAG CAGCTACCTA GOGGAAACAA GATGTAGTGC TATTGCCGAT AACAAGTAAG ATTTTCCACA CTGAGAAAGG AAAGCATTCG GATCTGCTGC AAAAACACAT ATATCCATAA AGACTCATGT TATTICAGAAA ACAGATTIGTIG AACACAATCA CATTICGCATG AATCCTTTAA AAGGAAGAAG GGTTATCTGA TTATTAGAGA TATTATTTTG GATATGTTAC TTATTAACTT GCTATGGCTG ACCITIABAGT ATCTGCAAAT CTGAATITICT AITTAITICCT TCACTGAATA TAGAAACAAT GTAACCATGA TAAAGTCTGT TATTAATAAC AACATAATTC TTTTTTTAAA GAAGAAAAGC TTAITITICA ITGACAGIGI AFAGAITITAT CTACITAGIT GIGITITIGCI ALTAGIGITI TAATTITITI ITTAAGITGA GIGITIGAIA AATTITAAGA CCCIGIGGCC ACCTIGITIT TAITTATAAA AGAATCMAMC MOTTGCATGC ATGAGGCTGT GAAGTCAGAT ATTTAGTAAT AAAAGCAGCA GIGCCITITIT ITGIATITIAC CCATIGACCC CCACCAAATG CAACIGITITI atattaagaa aatagtaaca attittaaaat ctcagagtaa aatctattitc actacatgcti TITCCCCCCT TGITCTGAIT TAAGCAGTGT GTACTTGGCA TCTCTACAIT GTCCTAGGGA CAGINGIGIT CTACAATAIT AICAIGTAIG AIGITITIAIT GGIGCITIITI AITICATAGIG GCTTCTTACC AGAACAGTA GGAAGAACA CATGAACTGT GTACAAGACA TGAAACATTG GAGTCCTIGTG TTGACTACAG GTATATAGCY CAMTITAAAA ATCCTAAAGC AAAAGAATTT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26: (A) LENGTH: 2036 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 26: CTAAAAAAA AAAAAAAA AAA

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900 960 1020

CTGCTGATAT GITGITITIT CACATGCTIT TGAGITITICA CITITITAAAC GAGAGCCAGC AAGCAAAATA GATGTGGCTG GGTCTGCCTG TCCGGGCGGC TYTYTGCACC GAGCTCTCAA

ATCCTOTOTA TIGAGGITC CITITIGGTA CICAGGATTG GAGCTACAGC TGGGCCCCCC TCTCTCCCAT TOGITTGAAG AGACACTGAG GGAAACAAGG GTTTCTTTTG AGGTGTCCTT

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CHOTTMOCIO G	CACCIONACCORC	360 GGCACTYSCTG	300 55		240 (xi)	180 50 (D)	(C) (B)	GOCACGAGAT AACATAGGCA CAATAATACT GTATGTCTAC TTCTAGGATT ATAAGGAATT 60 (A) LEGGTH: 556 bas	(xi) sequence description: seq id no: 27:	(i) SEQUENCE CHARACTERISTICS: (a) LENOTH: 717 base pairs (b) TYPE: nucleic acid (c) STRANDELWESS: double (d) TOPOLOGY: linear	(2) IMPORMATION FOR SEQ ID NO: 27:	35 ARGCTTCGAA GARGTAATAR AMCCCTGGAG A		2036	1980 30	1920	1860 25	TATTITITAA ATTIGICATA TATGGAAAGA GCATGTTIGT TACATGTAAA 1800	1740	1680 20 (D)		1560 15 (1) sego	1500 (2) INFORMATION 1500 15 (1) SEQU	1440 (2) INFORMATION 1500 15 (1) SEQ	1380 10 1440 1500 (2) INFORMATION 1560 15 (1) SEQ	1320 1380 1440 1500 1500	1260 1320 1380 10 1440 1500	1260 5 1260 1 1320 1 1380 10 1440 15	(xi) GAMTTCGOCA TATTTCCTTTT AACTGAACTG AGGCTTCGAA TTATTTTTTA ANGCTTCAAA ATGACTCAATC CACAGAAATC CACAGAAATC (1) (xi) AGCTTAACCT GACACCTCTT GACACCTCTCTT GACACCTCTCT GACACCTCTCTT GACACCTCTCTT GACACCTCTCTT GACACCTCTCTT GACACCTCTCTT GACACCTCTCTT GACACCTCTCT GACACCTCTCTT GACACCTCTCTT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACCTCT GACACTCT GACACCTCT GACACCTCT GACACCTCT GACA		CCTCCCTTAT TAMENTARIA TCTTCATGA GENERALLA ANDCETTCTT CCTCCCTTAT TAMENTARIA TCTTCATGTA TUTNTOCCH ANTOTAGCT GACTITIAM AGCITTICT TOTOTOCATG CCCTOTOCAG GENEROTAT TOTACATOCA TOCCTTTCOT CCCTOTTTCCT TOTACAGAT ANTOTAGAGA GENEROTAT TOTACATOCA TOCCTTTCOT AGCITTIACTG ATATACAGAT ATACTAATOT TICAAGATG TOTACTTCA AAGTOTACAG TITTCAAATG TIGTTACCAG TCAAACACCC TIGTGOTTA AACTTGCTAC AAGTOTACAG TITTCAAATG TIGTTACCAG TCAAACACCC TIGTGOTTA AACTTGCTAC AAGTOTACAG TITTCAAATG TIGTTACCAG TCAAACACCC TIGTGOTTA AACTTGCTAC AAGTOTACAT TAMTTCATT CCTCCCATGT AACTAAGAAT CATGACTATA TITTCATATCA ACGTTATATT (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 117 base pairs (B) TYPE: INLIEIC acid (C) SITRANDEENESS: double (D) TOPOLOGY: lineax (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 27: GOCACGAGAT AACATAGGCA CAATAATACT GTATGTCTAC TTCTAGGATT ATAAGGAATT AACATTGAG AGACCAGACT CGCACCAACA TTCAACCCCA GCGCTGATAT GAAAGTTTCC TCCCAGGCTA GAGCCAGAC CAATAATACT GTATGTCTAC TTCTAGGAAT AATATCTCCT CCTCAGGGCA GAGCCAGAC CAATAACACC ATCACAAAAA GTTACACAGAG TAATTCTCCT TCCCAGCTTC COGGAATCACC ACTTTCAGCTA ACCCAAAATAA CTCTCGAACT TAATTCTCCT CCTTTTACCT TTATCCATTTA GGTGAAGCAT TCCACAAAAA CTCTCGAACTT TCCCATTATA GGGCTGTGGT TCTCTCTGTGT CCTGGAATAAA ACCCAAATAA CTCTCGAACTT TCCCATTATA CATTTACCTT TTATCCATTTA GGTGAAGCAT TCCACAAAAA CTCTCCAACTT TCCCATTATA CATTTACCTT TTATCCATTTA GGTGAAGCAT TCCACAAAAA CTCTCCAACTT TCCCATTATA CATTTACCTT TTATCCATTTA GGTGAAGCAT TCCACAAAAA CTCTCCAACTT TCCCATTATA CATTTACCTT TTATCCATTTA GGTGAAGCAT TCCACAAAAA CTCTCCAACTT TCCCATTATA CATTTACCTT TCCCTTGTGT CCTGGAATAAAAAAAAAA
ç	TACTOROLLS DIALITY OF THE CONTROLLS OF T	ACTATOR CIPTOTICION GODACION TODOCANO CARACIDOTI CAO	ACCICIT AGAAGAGCIG CIAGAAAGGC AGACAGCACC AAGCGCIIAA AIG	TTAACGT CATGATTCAT TAGGGGAAATG CAAGGCAAAA CCATGATGAG AAT	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 29:			(1) SEQUENCE CHARACTERISTICS: (A) LENCTH: 556 base pairs	INF	AGAAATC CTAGG	ACTUAGT GGTGGCACAC TATGGAGTCC TGCCCACAAG TAGCACACAT CAA	CTTCGAA GARGTAATAR AMCCCTGGAG AGAGAAACTG AGACATGTAA GAG	TITITIA AAAAGGCICC ICCAGGAAAT GCATATAAGG GCTAATCACC CAG	rocogga tigaecetta caacatgeag togecetaca gaaaaacetg caa	SCATTIG CICCTACAAG CIGAAAGGCA CCCCIGGGIG GCIGGGGCCC ICG	IGAACTO TIOTTITCAT AGGINAANGA GAGACTGAGI TITITICATIT CIV	IOCITIT ICICCCACAA TIAATCIIGA TICIGCCIGI CIGIGCACAT TIG	ITCGGCA CGAGCAGCAT CCTAATTITA GITTGGAGAT GCATTCTAAA GGA	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:	(D) TOPOLOGY: linear	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 495 base pairs		INFORMATION FOR SEQ ID NO: 28:	INFORMATION FOR SEQ ID NO: 28:	INFORMATION FOR SEQ ID NO: 28:	NOTITGAT CAGACATTAG ANTANTTHATT OCTABABACTA AAAAAAATTA AAAA INFORMATION FOR SEQ ID NO: 28:	PAGATAG GGANOGTOGC GTANCTTCCCT ACAGITTCCC TGTIMACAAG AAAG GGTTGAT CAGACATTAG ATTATTTMTT GCTAAAACTA AAAAAAATTA AAA INFORMATION FOR SEQ ID NO: 28:	GETGATICTICA TACATICCTAA AGTITICAGAA CCATTGAGTA AAGTTAATICC ÅTTAAGAAGA GATTAGATAG GGATGGTGGC GTATCTTCCT ACAGTTTCCC TGTTAACAAG AAAGTCAGAG GTCAGTTGAT CAGACATTAG ATTATTTATT GCTAAAACTA AAAAAATTA AAAAAAA (2) INFORMATION FOR SEQ ID NO: 28:	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 495 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: Linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28: (TOCOCA CGAGCACAA TRANTCITGA TRUTCAGAT CCATTCIAAA GGARCTICTC TOCOCAA CGAGCACAA TRANTCITGA TRUTCAGAT CTOTOCACAT TUCCATGAGA TOCATTIT CCCCACAA TRANTCITGA TRUTCCCTGT CUTOTOCACAT TUCCATGAGA TOCATTITA CAACATGAAA GAGACTGAGT TUTTTCATTT CTGAAGAGAA TOCATTITA CAACATGAAA TACCCTGGAG AGAGAAACTG AGACATGAAC TOTOCAAA GARGTAATAR AMCCCTGGAG AGAGAAACTG AGACATGTAA GAGGGTGGGAG TUTTTTA AAAAGCCAC TAUGGAAAT GCATATAAAG GCTAATCACC CAGTATTTTG TOCOCGA TUCACCCTTA CAACATGAAA GCACTGAAC GAAAAAACCTG CAACTAAAAA CUTOTAA GARGTAATAR AMCCCTGGAG AGAGAAACTG AGACATGTAA GAGGGTGGGAA CCTTCAAA GARGTAATAR AMCCCTGGAG AGAGAAACTG AGACATGTAA GAGGGTGGGAA ACTCCAT GAGGCACAC TAUGGAGATC TACCCACAAG TAGCACACAT CAACCCACTA ACTCCATG (A) LENGTH: 556 base pairs (A) LENGTH: 556 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE CHARACTERISTICS: (A) LENGTH: 556 base pairs (B) TYPE: nucleic acid (C) STRANGENESS: double (D) TOPOLOGY: linear (xi) SEQUENCE CHARACTERITON: SEQ ID NO: 29: TUTAACGT CAUGATTCAT TACCCACATA CCATGATGAG AATGCCCCTA ACCTCTT AGAAGACTG CHAGAAACGC AGACGCACAA CCATGATGAG AATGCCCCTA ACCTCTT AGAAGACTG CTAGAAAAGC AGACGCACAC CAGCCATTC CTAGGTTACT TACCAAAGAA		20 20 30 30 35 50 50
	CAAGGAG	TCTOGGA	AGATOGG	GCCCCTA							CCCACTA	OGTOGGA	PATTITIG	CTAAAAA	TOGGAGT	AAGAGAA	CATGAGG	(ICITOIC								AAA	утськай	NGAAGA TCAGAG AAA		(1) GAMTTCGGCA TATTTGCTTTT AACTGAACTG AAGGCATTTGATTTTTAA AAGGCTTCGAA ATGACTCAAATC CACAGAAATC CACAGAAATC (2) INFORMA (1) (2) (2) (2) (2) (3) (2) (3) (3	20 (xi) 20 (xi) 23 GARITCOOCA 25 TATTOCTITT AACTOAACTO AGGOCATITG 30 TENTEUTYTA 35 ARGCTECAA TENTEUTYTA 36 CACAGAAATC (xi) 50 (xi) ACCTENACOT GACACCTCCTO GACACCTCCTT GGCACTOGTG GACACCTCTAACCTG GACACCTCGTG GACACCTCTAACCTCTAACCTG GACACCTCTAACCTG GACACCTCTAACCTCTAACCTG GACACCTCTAACCTG GACACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTCTAACCTG GACACCTCTAACCTCTAACCTAAC

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	. 257			258	
	ANTHGAAAGC AGGCAGATCT TTACAGCAGC TCTTACCTGH TTGCAAAACA ATGGAAATGC	300	CTGGCCTTAT TGGACTCCTT TTGGCTAGAG GTTCAAAAAT AAAGAACCTA GTGTATCCGC	AAAAT AAAGAAGCTA GTOTATCCGC	240
•	CCACATOTICC ACAAACAAT KTOTGOTICTG CCTGTGCCAT GAAGCACAG GTGGCTGAGC	360	CIGGITICAL GGANTIAGCI GCCICCCICI AITHICCACA ACAAGCCATC	CCACA ACAAGCCATC GTGTTTGCCC	300
S	OTCANGNOTO COCINCINCA ANGUISICAS CRISTACAGS GOTICACIACI OTGIGATTOC	420	AGGICAGIGG GGAGAGATTA TAFGACIGGS GFTTACGAGG ATATATAGIC ATAGAAGATT	CGAGG ATATATAGTC ATAGAAGATT	360
	ACACATOTICA CATTETTOGAC ACGGACATICE TOGATGGCAA AACGAGCATE GGGCTGAGAG		TOTGGAAGGA GAACTTTCAA AAGCCAGGAA ATGTGAAGAA TTCACCTGGA ACTAAGTAGA	AAGAA TICACCIGGA ACTAAGTAGA	420
01	GACTICCTGAG AAGGCGAACG GGCCTGCTGG GATGTGGGTT GATTGTAGCA GTAGCTCATG		10 AAACTCCATG CTCTGCCATC TIMATCAGTT ATAGGTAAAC ATTGGAACTC CATAGAATAA	TAAAC ATTCGAACTC CATAGAATAA	480
	GAGATOTGAC CTCAAA	556	ATCAGTATTT CTACAGAAAA ATGGCATAGA AGTCAGTATT GAATGTATTA AATTGGCTTT	GTATT GAATGTATTA AATTGGCTTT	540
!			CITCITCAGG AAAACTAGA CCACACCTCT GITATCITCT GIGAAATCAT CCTACAAGCA	CITCT GRAAATCAT CCTACAAGCA	600
15			AACTAACCTG GAATCCCTTC ACCTAGAGAT AATGTACAAG CCTTAGAACT CCTCATTCTC	ACAAG CCTTAGAACT CCTCATTCTC	099
	(2) INFORMATION FOR SEQ ID NO: 30:		ATGITICCIAT TTATICTACCT AATTAAAACC CAAGTTAAAA AAAAAAAAA AAAAA	FAAAA AAAAAAAAA AAAAA	71.5
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 414 base pairs		20		
;	(B) TYPE: nucleic acid				
	(C) SIYANDELMESS; double (D) TOPOLOGY: linear		(2) INFORMATION FOR SEQ ID NO: 32;		
25	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 30;		25 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 486 base naire		
	CTANATOCTO ACTOTGOCTY TOTCGAGACA GOCCCCANAT GOTNOSTOTG AACACAACAT	09	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	*	
2	GCACAGAATG AGGAGACATG CAGAGTGCTG AAATACTGTC CTGGACAGAT GTGTTACATG	120	(D) TOPOLOGY: linear		
3	ACTITICITIT CACCITATIT CIGIOCCIG CCITICAACA TAGAGCITIO TICATATIA	180	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 32:	ID NO: 32:	
	CATTANACA AMITGIATAA YIMIGIITCA TICIGACAIG TIMITIAGCA AMGAAAAR	240	GAGCCAGTGC CGGCGAAAGG GGACCTTCCT CTACTTCCTG CCACAGACCC TGTCCCCACA	CCTG CCACAGACCC TGTCCCCACA	09
35	GAGTARTICT ACATCAGCAT CTITIAGISCA TGCHAAAGA TITAAAAIGT CTITITISGGA	35	CACTICCTISC COCTIGCTOTS CTIGGGAGGCC ACTICCTICCC CCAGTGCTGG ATTCCACCCC	TCCC CCAGTGCTGG ATTCCACCCC	120
	ACAUGITITIG TATACATAAA TGTTTAGATA GAAATATTTA TAGAATACTC TATGTGAGTA	360	CAGCICACCC ICAAACAIGG CCCCCICICI CCTCCTGCII GCCCCTCTCI GCTCCCTGGA	GOTT GCCCCTCTCT GCTCCCTGGA	180
9	THUNICICCC INTOININI TAINICING TOTOTCANIC TITGINING THIGHANICC	420	GOCTETTCTG TOCTCCCTC TTGAAAAGCA ATGCCAGCTT CCTGGGATCT TCTGCCAACT	GCTT CCTGGGATCT TCTGCCAACT	240
9	tatgaatagt gaga	434	CONGCTACCA TOCCOTTIGG TOCTOTCAGE TCAGGETCCTC AAGGGAATTG TCTAMACTICG	CCTC AAGGGAATTG TCTAMCCTCG	300
			GIGTECTIGCT TECCTECTE AACTECTICA COCTGCTECA AGCTEGCATE	TCCA AGCTGGCATC TGCCCCTCCA	360
. 45		45	CTGCACAGAA COGNTCCCCC ACCACCTGCC TTTACAGGGA GGAAGCAGCA ACATGGAAGA	GGGA GGAAGCAGCA ACATGGAAGA	420
	(2) INFORMATION FOR SEQ ID NO: 31:		ANCIBARCTAT AGGGGCTACA ANGATGCTCA GCTCTGATCC CGAAGGCAAA AAGINTCTTT	ATCC CGAAGGCAAA AAGNATCTTT	480
80	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 715 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: double (D) TOPOLOGY: Linear	50	GGGCAC		486
Ş	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:	55	(2) INFORMATION FOR SEQ ID NO: 33:		
3	CCACGCGTCC GATCTCACAG CTCCGACACT ATTGCGAGCC ATACACÁACC TGGTGTCAGG	09	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 725 base pairs	g	
	AAACGTACTC CCAAACTAAG CCCAAGATGC AAAGTTTGGT TCAATGGGGG TTAGACAGCT		(8) (8)		
99	ATGACTATET CCANATIGEA CETECTIGGAT TITTTECEGAG ACTIGGTBIT APPOSITITIG	180	(0)		

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(2) INFORMATION FOR SEQ ID NO: 35:

	AAAAAAAAA AACTCNA	CTACATCTICC AGCTTGAGGT TROCCTCATA TCACATTRICA TTCTCACTAN ARACHAMANA	ATCACCTOOC GTOCCCAGAT CCTCOCARGO CAACACCCTG TGATHAATTCC AGGTGATTCT	AGGAATCTTA THACCTACGT GGACTCTTTC CATCCGTACA TEGICGTOCA CATGCCACTC	CCAACGTOGA AAAGTATTCC AGGTCCATCC CCAAGGAACC AACACGGATG ACATGGACTC	CTCCTCCTTT THTTCGCTCT TGTCGAAATC AAATTGGAAG ATCTTCAGTC CCAGCTGCAC	TOCAMBATAC TEMBAGATTO AMBCATOCTC TOGAMATOTT CCCAGMACAT TOCACGATGC	CACACAGCAT GCTGCCCTCA GACGTGTCCA TCCTGTACCA CATGAAAAGG CTGCTGCTCC	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 34:	(D) TOPOLOGY: linear	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 437 base pairs	(2) INFORMATION FOR SEQ ID NO: 34:			АААА	CTTCATACTT TYCAATTYGA TAGAAATAAA CHTTTTTTCT CCTTATAAAA AAAAAAAAA	TGIATATOCT AUCCIAACIG TIAATIGIAT TATTGATIAT GTIGATIATC TIGCTIGAAG	TIGGIGGCCC ICTIAATTIT GGIGTATGIG CTICCAAGTA ICTAAACCIC CAGICIGAIC	CAAGITITCCC AGAAGICGIG IGITIAIGAI GAGICAGAGI GCITITICCIC GGIGGGAGAG	CAGITICIATA ACCCAATGAC AACCTOTOTO TITTOGITTIAC TOTOCTOTGA AATGTCAGCT	TOTOTOGOGA ARTIGITIGAG TIACAATOGC ATTICACTOT GATOCCTOTO AAGCTCAGAT	TOTAGTITTA TGAGAATITG TACTACTGAT TITTATATAT TCCTGTTTTT GATGAACAGA	AGANANATIGA ANACTICAGTIC TITTATISTANG CTOCHAGGAT ATTAGGGCTT ANAGGGCTTT	AGTCAGAAAT GTCCTAAATA ACAAACTATT TTGTATTAA TTTAGGGAAG ACTAAAGGGA	AUGIGICIAA TICATITTAT AAAAATIKIT CIUGICIICA TITIAAAGU TUGGUKIKU	TTTCCTGTTT AAAGCTTTTC AAAGGAGCAG ACCACCTTGA AGATTCCCCC TAGGGTTGAT	GTTCCTCTGG TAATAATTAG GTTATTCCCA GAAGCACAGT GTCATTCTTT AAATAAAAGC
	437	420	360	300	240	180	120	60								725	720	660	600	540	480	420	360	300	240	180 .	120	60
60		55			50		45			40 .		 			30 ,		2		.3	20 ,	د.		15	6	10			5
GAATCCTAIG TCTCGCCTGC AGGTGGTTGG TTTTCAATOT TCTTGCTAAT TTTTTTTCTA	GECACGAGAA ATETTEATICE TOTAGTEACT CEAGACEATIG GAGTOGETIT CEAGETGAAT	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 36:	(D) TOPOLOGY: linear	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 604 base pairs	(2) INFORMATION FOR SEQ ID NO: 36:			Малалала Лалалалал Лалалалала Лалалалал Ала	AGTGANTTGT ATGTGGTTAA TTATAAATAA AACTGGTACC AGGNAAAAAA AAAAAAAAAN	TITOGITITIC TCATATITNI GIGNATIAAG GNAIAGAIGI TAACCNITAT TITGIAGNA	NOTICATINIC AACCIVITGIA TIVAARGCITA GACTAAATAG TAATATATING TOGGIAGGAT	CAGATTACCC ATNTOCAGAA CTAAGGGAAG CNATTTATGT ATGAAAGNAA TTNTTGAATT	ACTAINTIGT TGATAOCTIT TIGTICININ AGGITGUAAN ATGACAGIGC INAINTGAAA	TOCACTIAAT ACGCACCIAT TINTCAATAG TOTTATITIT TOGNIAGCAT TITTITIACC	TOTICCTITIAG TOCCCTAAGG CTAAATTITIG GTCATTIGAC ATCAGAGAIG TIGIAAGTAI	THANCHICAC TITHAGIGAA AHAAAAIGIG CCAHACHAGI AIGIGCITCA AAAGGGCAAA	AMACANTOGT TTTTTTNGCC ATTTGACTOG CTCTTTNAAT AGTCTACAAG ACATTCACGT	TTGAGTGAAA TATCATAAGA TGANAATGGA AANAAGGAGA CACAAANAGT TATNACAAAA	ACHITICITA CACCCIGGCA GAAGGGAGAG AAAIGIGITI IGGGGIGGGI AACIAAAIII	TCTTTGTTCA TTANCATGCT ANTICCTCTT CAGTGTTAR TTTCTAGTGA CAGAATGCTA	GANTGACCAG AUGCTIANGG TCIACATITT CCITTANCCT GITAGIATIA CCITCCITAA	AGAAGTACTT ACCICTIGAA GATITAATAT ATAATGGTIG ACATGATACA TGTACATGAT	GOCACGAGCT GGAACAGAGA CTAAATCCCA CGAAACTGAC ATTOTTAAAC ACACTAAAAC	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 35:	(D) TOPOLOGY: linear	(B) TYPE: nucleic acid (C) STRANDELWESS: double	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 943 base pairs
120	ě	;							943	900	840	780	720	660	600	540	480	420	360	300	240	180	120	. 60				

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

M	WO 98/39448 PCT/U		WO 98/39448	262	PCT/US98/04493
	261			707	
	TIGGATCTIG GGAGTTTTCT TIGTTTGCTC CTGTGTTTGC CCAGCTTTAA TAAAACCAGG	180	GGAAC	GGNACGGCNG CNGCGGTCNC NGGCNAGTNA NTNGTNATIGC CGGNGCNAGT TTCCTCCGGC	180
	COCAMACAMA AACCATROCA TTCTGAACAA TAGGGGGCCC ACATTGGACC CAGTATGTCA	240	TTTAT	TTIATCATET CACCCACTET GGTATATGCG TTGTGGTCTG CCAACTTTGC CGTGAACAAT	240
S	CTTTAARGGA CTTCTAGAAA AAARCTGAAF GOGAAAAATG ACACTAGGAA TGTARACTCC	300	5 TTCAG	TTCAGCAATA ATCAGATGGC GGCTGGCGCA ATATTCAAGA TAACGCCTGG CAGTGGTGGG	300
	ACACATITIA TOCCATATAA TOGTOTOTIT TOTTAATITIT GITTCTTGTG GCGAAATGTG	360	GCTGA	GCTGATGGTT CAGTGCCTGC GSCACCGTTT YTGCCGTATG TTGCACACCA GGNTCTTTAA	360
	SCITICAAAT TAAAATGACC TITICTICIT TGAAACTTIT TGTITIGACT TGTATAATTA	420	ACAGT	ACACITITICS SACCECCITT ACCOPCAAGG GITCAANGCC GGTCGGIAGC TCGTCCTTAG	420
0	AGGOTTICGA AAGATICATA ATTCTGAGAG AGGITTICCAA CCAGGAGATA CAAAGAAGTC	480		GITCACCGCG ASCATAAGCA ITAAACATCI CATCAATITG CITCTGGCTG GCGCTATCAA	480
	TCAGTAGTAA TCTTGTTCAT GTGCTTTTAC AGCCAGCTAC ATTTAAGGAT GTATTAGTTA	540	TACTT	TACTITICCAG CATATGITIA CGCTGGGGGA AACGGGTTAG CGTTTGCCCC ARCAGATCAT	540
15	CAGAAATTAT ATGTCTGTG ATGTGTCTCT ACTCAATAAA GTACATGCCT CCACAAAAA	900	15 лосся	AGGCAATGGG CTIVATGAGA IVATCAAATA CACCACAAGG TAGGGCTTCA GACACGGTTT	009
	АААА	604	CCATA	CONTATOBET GECTGCAGTS STAMACACCA COTCOCCOGG ATANTOCCCC TGCACCAGTT	099
20			carec	CATGCAGTAA AT	672
	(2) INFORMATION FOR SEQ ID NO: 37:				
٧,	(i) SEQUENCE CHARACTERISTICS:		(2) I	(2) INFORMATION FOR SEQ ID NO: 39:	
3	(A) LOWGHT : 47 DATE PAILS (B) TYPE: nucleic acid (C) STRANDEONESS: double (D) TOPOLOGY: linear			(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1908 base pairs (B) TYPE: nucleif acid (C) STRANDEDRESS: double	
30	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 37:		30	(D) TOPOLOGY: linear	
	GTGAGTGCCC GGGAGCCCG AGGCCTGCC CCTAAGAAGG ATATCTYTFA CCGCTCCCTT	09		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:	
;	TICACACACC TAACCCCCA GETGETCAGG CAGTGGGCAC ATGGCAGGGG CETCACTGGG	120	35 AGAGT	AGAGINGAIA ITITIAGAAA CAOTAAITII ACITIIAAGO AAAITGGCIA GCICIIIIGAC	09
32	GOCACATAGA GCATITGGGG GACTGCGAGT GCTCACCTTT GACTTCCTGC AGGTCGGGGG	180	_	TRUBGAGCIG TAGGAAGCIC AACAITICIT TOTAGAGAAC GITGCITITI TIGGAITIGTA	120
	AAAACCAGAT CAIGAIGACC AAAGTYIACA TATTCTIGAT CITCAIGGIG CIGAICCIGC	240	CAGGT	CAGGIATAAA AACATTGCTT TTGTTGAATT GTATAGGTGT AAAAAGGGAA TAACTGTATG	180
9	CCTCCCTGGG TCTCACCAGG TATATGCCAC CACTTTCTGY TCTAAATTCA GAATAAGAGT	300	40 слост	CAGGITTGAA AAGGAAATGI GCTTTAGGCA TGAGTCATAA GATGCCATTG TACTTGTAGG	240
	CACATCAGGA GAGCACTOTO COCAGGANAA TOCAAAOGGG TTOGCAGCA	349	CATT	CATITIAITI TOCITIAGAA ATGGACATCA GCICTICTCT TCTGACTGGT AACACATAGC	300
			CCCMA	CCCAAAGCAT GAGATTATTT TTCATTGGGT TTTTATTGTT GTTTAGTTTT GGTTTGTTAC	360
. 45				GCCAGGCCAG TCTGTCTGCG GAACACTGAC TCTGCTCTCT AATGAGAACA AAGTTAGAAA	420
	(2) INFORMATION FOR SEQ ID NO: 38:		TCTGC	TCTGCCGATA ACCTAAAATA ATTTAGAAAT GAATTAAAAA TGTGAAATGG GGTTAAAGTG	480
20	(i) Sequence characteristics: (a) Lensth: 672 base pairs		50 ATCAT	ATGATGATAA AATMGCATGC AAGAAACAAG CTCCTTCCAT CAGACTTGGC TACTGTTTTC	240
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		TYCY	TICHGGIAGG ATTIGGTITG GAAGAGCCTC TIGTITCCTT CTCTTTGGGG TAIGTCTTCG	900
				TITICITIALIA TOTITICIAAC ATTATIGAGA TATAATICAC ATACCTIACA ATTICACTIAT	660
55	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 18:		CC AATTT	TITIANGGETA CARITITAGIS GITTITAGIS TATTCACADA GITGISTAAC COTGACCACA	720
	GTAGTOSTIG CGSTIGCCGG GATGGCGAAG ATCTCGCCGT TIGAAGTCGT AAAACGCACC	09	GTCAA	GICANTITIA GAICAITITCG TIACCCCAIA NAGAAICCCT GIACCCTICA GCAGICACCT	780
99	TOSTIACOS TECHTOTISS TITISTISMIT STANTOTIS CTACAGACT GATGSTCCA	120	60 стсм	CTCATTITICT CCCAGTGCCC ACCCATCCC CGAGCCCCKG GAACCACTAA TCTATTICTC	840

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;	ааа		55 COC	TT	50 TTG	באניד		45 ATC	AACC	40 cna	CAGC		35 000	ATCC	30 cnac	TTAC		25 CTG1	930	20 ages		15	10 (2)	5 CATA	
	AAAAAAAAA AAA	САТСАСТТТС СССТСАЛАЛА АЛЛЛАЛАЛАЛ АЛЛЛАЛАЛА АЛЛАЛАЛАЛА АЛЛАЛАЛАЛА	CCCCCTTCTT TCTCCACTGT ACAGAAGAGC CACCACTGGG ATGGGGAATA AAGTTGAGAA	TITAANCTITI GCTGACGOTT CAGTCCTGCC TCTACTGTCT CTCCATAGCC CTGGTGGGGT	THOGAAGGG CAGCCCCCC NGCINCING GITINGIGGT NGCCAGCCIC AGGICAICCI	THOGGAATHO CACTITHOGG CCTTHOGGCT CHOGAACCTO CHCHOGGTCA THOGHGAGAC	croccerero adcoresero gareneroso adencertoa todenecead acertoserr	ATCACTGATG COCTGGGCAC CTCAGTCACC ACCACCATGC GCAGGCTCAT CAAAGACACC	AACGTGTTTG ACACAGGCTT GOGGAACGTG CAGCCCTACC TGTACAAGAT CTCCTTCCAG	GTGGAGGTCA TGAATCOOCA TGAGTACTTG CCCAAGATOC CCACACAGTC GGAGGTGGAT	аносооська тепосоосая самоопоост садаластет оссыдалалт сапсалсате	TOCATICCCAT GODAAGTOTO GACOGTICAAG GTOCATOTOG TAGCCCTOGC CACOGAGCAG	ососнантот осттоанотт стиссивана ананнотого остоосситт стонвновна	ATCCGTGCTC CAAACTCTAC ACTCAAGGAT GCACTGCGCA ACTCTGGTGG CGATGGGCTG	стосодност соснасьного сотоновного навинанного оснтаноска насттоновн	TTACGIOGAC IGIOGAGCIG GICICTIGIG GCICAGCOCC GIGCOGAGGI IGAAGCGIAC	TOCHACCHAC CTOGGCGCGG GGCGCGCCC GAGAGACCCG AGGAGTCCGT TCCTCCCTGG	спотоссоос отваваниес сессоована тавинесена осветестве насттесные	gacragreer galererger resessalager gaglacrare tellecterer geoecetere	GOCACAGAGC CICCGACCCA GOTOOTCIOG AGCCTGOCGG GAGAGTGGTG GCATCTGAGA	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 41:	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1153 base pairs (B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLOGY: linear	INFORMATION FOR SEQ ID NO: 41:	CATAAAACCT CATAITITAA AINAAGIIGA AATTIGAA	
	1153	1140	1080	1020	960	900	840	780	720	660	600	540	480	420	360	300	240	180	120	60				458	

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GCCATAAAGA GAAACAAAAG ACAATGATGG TATTCTCTGT GTCCTCAGCT TTGGCACTTT

TAGCTMACCT GGGGAGGAAA TGAAAATTTC CTTTGTGGAT CTCCCCAAAT CCATTGTTGT TOTTGATGTT GCTAAGGAGC AGTGACCTTG CTAAAAAGAC TGAATAATCC ACCCACTGAA

300 240 180 120 60

8

CCTCAAAAAA AAAAANGAAA GGAAAGAGGT CTCTACACAA GCCCGTGATT CTTCATGGCA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 458 base pairs

AGGGATAACA TCAGAAATOT TTCATTTYCK GCTATTAGTT TCCAITCCTT TCCCCATCCA

6

(2) INFORMATION FOR SEQ ID NO: 40:

(1) SEQUENCE CHARACTERISTICS:

35

TOTTAAAAAA AAAAAAAAATT AAAAAAAACTG GGNGGGGGGC CCGGTACN

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TCTCAGTGGC TTGACAGCAT CTTCCTGGTT GTATGTGGCC TGTTTACATG ATGTATTGAA

TGAGTTGGAG GTGATTATTC TCTGTAACTC CCTAATGATT GTTTTCTAAG CATTGTGGCT ACTCACTTAA TAACCAGGGA ACCAGCCAMA TACTGTGCAG CCGCAGAATA TGCATATCAA GACTGATTTC AGTGATATTC AGAAGTGTGT ACCAATCAAG GCTCTTTAAA ATACGGAACG CAGTCAATTA TICCCTAGGG TAGTICAAAA ATAIGAIGIG AGCTAGTTAA OCCITIOCIT

1680 1620 1560 1500

TAATGITGIT TOTIGIGAGC AICAATGCCT GTAACACCAA ACTAAACACG TGITTITTGGG

ATATOTTICC AATCTTTAAA TGACCTTOCC CTOTCCAATA AATAAATGAT TGTCTCACCC

1860

1800

1908

25

20

GATGCGTTGT AACACTGCTA AATATGCTAA GTACAGAATT TTATCTACAG TACTGTGAGA

1440

1380 1320 1260

ACTOCAGIGA GITTICIGCIG TOTITITICAA GIAIGIACCA TAGGACITAA AGGIGATITIG

CITTIOCTAGE AGETTOTATA CCTCAGGCCA GGTGAGCTCC CCAAATTTCT TTTTTCATTT CAMAMATCAG GCCAMATGAC TIGGCAMATA ATIGACAMAG IGGITITICAC GIGIGICTAT

15

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ACCAAGCATT TATGATATAT TACTACTTAT AATACTGTGG CTAGTCTCTA GAATGGATGT GGGCCAGATA CCTAACAGGT TTTTCTCCGT GAATCTTATG CTGAGTAGTT TTTCCTCATA TGAGAACTCT TATTGCTGTG AAAGGGAGTG GTTGGTAAAA TCAATAGATT TCAGGCAAGA

1080

1020 960 900

TGAAATCTTT GCCTCCTCAG TCCCGAAGAG TCCTGCTAAA AATCAGGCTA AAAATCAGGC

1200

S

TITITIGICACT GECTICCCAA ATAIGATITT CTATAIGGAG IGAGAAAATI CITICICAICT TCTCTGTAGA TTTGCTTATT CTGGTCATTT CATATAAATG GAATTCTACA ATATTCGGTC

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3)d SPP01/80 U/N	CTATIS98/04493	WO	WO 98/39448 PCT/A	T/US9	PCT/US98/04493
•	265					
			,			
			-	TAACTCATCC TAAGTGGGCA CATTTAGACA TAGCAGGGGT GATGACCAAC AAAGATGAAG	1500	
	(2) INFORMATION FOR SEQ ID NO: 42:		•	TTCCCTATCT ACCOLANGOC ATCACTGGGA GOCCCACAAG GACTCTCATT CAGTTCTTAC	1560	
S	CHARACTERISTICS:		2	TICGITICAG ICAAGACAAT GCTIAGITCA GATACTCAAA AATGICTICA CICTGICTIA	1620	
			~	ANTICCACAG TICAACITAA AAGOFFIFIG AATAAATOGA TGAAAATCTF TTAACGGAGA	1680	
	(C) STRANDEZNESS: double (D) TOPOLOGY: linear			CAMOGNIGG TRITTAMAA TGIAGAACAC ANTGAARITTI GINIGOCITIO ATTITITIT	1740	
9	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 42:		2	CATITICACAC AAAGATIAT AAAGGIAAAG TIAATATCIT ACTIGATAAG GATITITIAAG	1800	
	OCCUCANDA GOOCCANDCC GACAMANTOT TCTTOCTOCC TCTTCCGGCT GCGGGGGAA	09	•	ATACTICTATA AATGATTAAA ATTTTTAGAA CTTCCTAATC ACTTTTCAGA GTATATGTTT	1860	
15	TAGTOGICOS ACSTETOSCO OTGAGACOTT TOGGGAGOCOS GAGTOTOTOC ACCOCAGACA	120	15 1	ttcattgaga agcamattg taactcagat ttgtgatgct aggaacatga gcaaactgaa 🐪 🔢	1920	
	TRACGAMOGO CETTOTITTA GOMPICTATT CCAMAGAMA AGMAGATGAT OTOCCACAGT	180	•	arthctatg cacttgegg aaacaataaa tgcaacttgf tgtgcaaaaa aaaaaaaaaa . 15	1980	
é	TCACAAGTIGC AGGAGAGAT ITTGATAAAT TGTTAGCTGG AAAGCTGAGA GAGACTTTGA	240	* (c	AAA	1983	
₹	ACAITATETICS ACCACCTETIG AAGGCAGGGA AGACTEGAAC CTITTIATIGGT CTGCATCAGG	300	3			
	ACTITICCICAG CGTGGTGCTA GTTGGCCTCG GCAAAAAGGC AGCTGGAATC GACGAACAGG	360		ון אופריפוראוידראו מיס במי דו אול. או.	··· <u> </u>	
25	AAAACTGGCA TGAAGGCAAA GAAAACATCA GACCTCCTGT TGCAGCGGGG TGCAGGCAGA	420	52			
	TTCAMENCET GENERATED TETSTINGENED TECHNICOCTS TEGNICACET CAGGETTECTS	480				
ç	COGROGGAGE GGTOCTTOGT CTCTATGANT ACGATGACCT AAAGCAAAAA AAGAAGATGG	540	Ş	(c) STRANDEDIES GOADIE		
ş	CTGTGTGGGC AAAGCTCTAT GAAAGTGGGG ATCAGGAGGC CTGGCAAAA GAAGTCCTGT	009	3			
	TIGETICIDO GEAGAACTIO GEACOCAAT TGATGGAGAC GECAGOCIAAT GAGATGAGG	099	£	ANTARCANCA CHARLES CANTERLACT CHARLES AN ACCURACY CHARLESTERS	Ş	
35	CAACCAGAIT TOCCGAAATT ATTGAGAAGA ATCTCAAAAG TGCTAGTAGT AAAACCGAGG	720	35	AAGACHTIAF AGARATRGAC ATTTCTTCAA GGAGAAGAGA AGATCAGTCT	120	
	TOCATATCAG ACCCAAGTCT TOGATTGAGG AACAGGCAAT GGGATCATTC CTCAGTGTGG	780	. t		8	
9	CCAAAGGATC TGACGAGCCC CCAGTCTTCT TGGAAATTCA CTACAAAGGC AGCCCCAATG	840	• •		2 5	
5	CAAAGAACC ACCCCTGGTG TTTGTTGGGA AAGAATTAC CTTTGACAGT GGTGGTATCT	006			9 9	
	CCATCAAGGC TTCTGCAAAT ATGGACCTCA TGAGGCCTGA CATGGGAGGA GCTGCAACTA	096		-	§ §	
45	TATISCTICAGE CATESTICT GETGEAAAGE TTAATTTTGCE CATTAATATT ATAGSTETISS	1020	2 5	CIGATITIONG CAARITITIA TOOPACITITI TAAATAAGCCI TOTTACONGC AATICNGAOT TOOPAGAATAAA COTOTOTOTOTA AAAAABAAGG CITAAAGAAA ATICTIGAAG CAAAAAAAAAAA	360	
	сосстетта талалата сеслосваса деоселлела оссовават оттеттивла	1080	٠ .	·	9	
ç	CCAAAAACGG GAAGACCATC CAGGTTGATA ACACTGATGC TGAGGGGAGG CTCATACTGG	1140	. 05	CITIANNIA IGACIAACA AFAATITAAA ACAAICATA GIACACCAT	240	
3	CTGATGCGCT CTGTTAGGCA CACAGGTTTA ACCCGAAGNT CATCCTCAAT GCGGCCACCT	1200			9	
	TAACAGGIGC, CATGGATGTA GCTTTGGGAT CAGGTGCCAC TGGGGTCTTT ACCAATTCAT	1260			9	
55	CCHOSCIPTIG GAACAAACTC TITCGAGGCCA GCATTGAAAC AGGGGACCGT GTCTGGAGGA	1320	55		9 5	٠
	TOCCICICIT CGAACATTAT ACAAGACAGG TTGTAGATTG CCAGCTTGCT GATGTTAACA	1380	•	CITALITATION OF THE CONTRACT TH		
(ACATTOGANA ATACAGÂTET GCAGGAGCAT GTACAGETICE AGCATTECTG AAAGAATTEG	1440	, ,		8 8	
9				GIRGCAGIST GIGGGALGAG GITCATACA GACGFATTIA TIOCTIGICA IGIAAATTAA	840	

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8 6 ႘ 30 20 3 25 2 55 5 S CATGAGTCAC AGCITTIGIT CIGIOGIAAC CIATAAAAAA AGIITGICIT TGAGATTCAA CARGARAGAC CTGGATGACA TCAGCACCAA AACAGGCATC ACCCTCAAGA GCTGCCGGAG ACTOCACCAG CTCATCTTCC AGATTCCGCC CTCCCGGCAG GCACTACTCA TCGAGAGGTA CATGGACCAT TACCGCACCT TCCACATGCT CGAGCGGCTG CTGCATGCGC CGCCCAAGCT GOTICCOCTICG GGAATICCTIGG AGCAGACTIGG COCCACGGCA GCGGTISCTIGC AGAGCGACAC CGAGGACGTG TATCGCCTCT GGCTCGATGG TTACTCGGTG ACCGACGCGG TGGCCCTGCG CCCOGGEOTC GCCATGACCA GTGAGCTGGA CATCTTCGTG GGGAACACGA CCCTTATCGA GOGCCTGAAG GCGGCRCGCC AGTCCCGAGC AGTGCTCGCT CCTGCTCGGG GCGCTGCGGC 2 GCCCGATTAA NTGGTTTGAA GNCTTG TAGTTATGIA GITATTATGA AACCACCAAG ATTITTITTIGG CTATTTACCG TAACCAAAGG TAAAATAGGA AATTATAAAT ATATAGTTTT AAGCCTGCAT CAGTGGGAGT CTTGGCTATG TCATAAGAAA TTAAAAGAAC TTACCAGGAA GGTTTTTAAG TTAGAAATAT TCCATGCCAA TOTAMAGAAC TGAMAACMAT GTATATOTTO TAAMTATTTO TOTOTTOTGA GAMATTTTTTO CTACAAGGAA TATTAAAAAA ATCTATTCAC TTTAACTTAT AATAGTTTAT GAAATAAAAA CTAATGAACT ACAGCTATCT TAATTITOOTT CTTCAAGTTT TCTGKTGCAC TTGTAAAATG TOGTOCTTAT TITITICAAAA ATTIGCTOTG AACAACGIGA IGACAACAAG CAACATITAT AAACCIIGIA TITAACICIT TICAATCCIT TIAGATAAAA IIGIICIIIG CAAGAAIGAI COTOTTOTT GOTAACAACO GOTTTGAGAC AGGGAAGAAA AAACTGCAGT ATOTGAGOTT GGACAATATT CAGCAACACT TECTECTETE TGACEGGTTG GECAGGGACT ATGCAGCEAT CTATISCETTY GATGAGGEET TIGTTEGGGA GGTGETIGGGE AAGAAGETIGT CEAAAGGEAE CTCACAGATG GATGACATGG ACATGGACTT AGACAGGAAT TTCTCCAGGA CTTGAAGGAG ACAGITITGAC AACITITAAAC OGGICTICAA GGIGGIAGAG GAAATGCGGG GCICCCIGGI COGNIGACITY OCCINCIBES CHBASCICAY GANCCAAAAC IOGACCCTIG GACCCOICGA INFORMATION FOR SEQ ID NO: 44: £ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 44: (A) LENGTH: 1391 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 1080 1320 1260 1200 1020 780 660 600 540 480 420 360 300 180 120 960 900 မ 25 20 50 3 8 33 5 10 S AGCCCAGACC CACTCCCTOC TCCAGCACCA GCCCCTCCAG CTGACCACCC TOTTGGACCA

GOTTCTTCGCT CCGGGGAGCA AGTOGGGGGC GTGCAGATIGT GCCTGTGTCT GTCTCTGAGC OCCOTTETTE CTGAATCAGT ATTEAGEGTE TOTECAATEC CTCGATGGET TECGACACCA GGGGGCTGGT GAACGTGCCG CCAAGCTGAC CCACAATAAA GATGTCAGAG ACCTGTTTTGT CICAMOGICC TAGIOOCIGA CAMOGACCITY CICCACCIGC ACAMGAGCCIT GGTGTGCACT CIGAGOIGCC TCCCAACGIC COCCCACGCT GACAATAAAG TIGCICIGAG TITTGGAGACT GOCCETETGG GACCOCTACA TOGOCACCCT CCGCGGCTGC CTCCTGCGCC TGTATCATGA GUACCITOSIG GAGAAGIITIG TOGAACCCIG CCGCITCOGAC CACIBGCCAC TCAGCGACGI GCTCTCCGGG AAAGCTGGGC GTCTTCTCTG AGATGGAAGC CAACTTCAAG AACCTGTCCC CICAGGGAAI GGIGICIGIG CICAGCCCAI CCACCAGAAG AGICIGCICA CAAAAAAAAA алалалала а ACCIOGIOTO COTOTACAAG GATOGATOTO INCHOTOGOCI CCTTGGGAAC TGAGACATAT 1380 1140 1320 1200 1080 1020 1391 960 900 840

3 INFORMATION FOR SEQ ID NO: 45:

E SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1569 base pairs

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:

COGTOGTOTO ATGUACTITIC AGCOGLAGAA GCTGATGGCT GTGACTGAAT ATATCCCCCC GECACGAGTG GAGATGOCTG CGGCCGTGGC GGGGATGCTG CGAGGGGGTC TCCTGCCCCA GAGGATTICCA AGTACCAMAA TCTGCTGCCC CTTTTTTGTGG GGCACAACAT GCTGCTGGTC GANACCAGOC ATCCACCCAT CATOCCTGCC ATCTCCTCCC AGCCCCCCAC GACGGGCCGG CIGCCTACCC ICCAGACIGI CCGCIATGGC ICCAAGGCIG ITACCCGCCA CAAGCTCCCC AGCCTGCCCC TGGTGCAGGG GGAGCTTGTA GGAGGCCTCA CCTGCCTCAC GCTGCTAGGT GGCTGCATTG ATGACACCAT CCTCAGCAGG CAGGGCTTTA TCAACTACTC AGTGAAGAGC CCAAGGTCAA GGAGATGGTA CGGATCTTAA GGGACTGTGC CATTCCTGCC TOCOGRADACA CARGATECTO ATGARÁGICT TECECLARCER GOTECTORAR GECETTÉCTO ACCCONCIGE CAGAATOTOG CICTOAGTOC AGAGGACAAG CITICITATIG AGGCCTCATC AGGCTTCTCC GCCGGGAGAT AGCAGCAGTT TTCCAGGACA CGACACCAGC ACCGAATGAT AGGAGGAGAT 540 660 600 480 420 360 300 240 180 120

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GTACATCAGA GAGCAACGCG AGRAAGGATT CTGTCATGTC GGCCAATGGG AAGCCAGATC

3	WO 98/39448 PC	PCT/US98/04493 W	WO 98/39448 P
	269		. 270
			CCCAAATAT ACTATCCCTT ATGTGAAGGT ATGTGACAAC GTTGACCTCA CCAAATGAGT
	CTGACACTOT TOCOGACTOG TAGCCAGCOT GTTTAGCCAG CCCTGCOCAT AAATACACTO	840	THIMACAICA GCICHTITT CAIMIGNANG CACAIMCCCT GCICCCCAIT CANGINIGIC
V	TOCSTIATITG GCTGTGCTCT CCTCAATGGG ACATGTGGAA GAACTTGGGG TCGGGGAGTG	900	TTCCATTGTC AGGCAGGCTG ACCACCTTCA GCAGGAGTCC TCCAAGAGTG CCCAACTCCC
,	TOTITICICAC TTGGTTTTCA CTAGNANGA TATTGTCAGG TATAGGGCCA CTTGGAGATG	096	CTTCCCACAG TACACAAGGC TGTAGTTGTT GTCCTGCAAT CCTTTGTATT TACCTCATTC
	CAGAGGATTC CATTTCAGAT GTCAGTCAGC GGCTTCGTCC TAAGTTTTCC CAACTTGGGA	1020	ITTCCCARCT ANGTECTICAE TGAGTITITAA AGITAGGGCT GGAAAAGCTA TGCCTTACTG
10	COTGATAGGA GCAAAGTCTC TCCATTCTCC AGGTCCAAGG CAGAGATCCT GAAAAGATAG	10	GGACAGCAAG GAACCAATTT TTTTCTGAGG GAGAAGACAT TCACCTTCAC TATATGCCTG
	GOCTATITOTIC COCTIGOCOTICC TITOGICACTIG COCTIGOTIC CACGOGOTICC TGAGGCOLACC	1140	GCAGGGCCAC AGTGCACAAA ACAAAGATCA GCCTTCATTC AAGTTCCAGG TTTTTTCTTCC
7	CCCTTGGGGC ACAGCTGCC ACTGCCACAG TAGCTCAACC AAGCAGTTGT GCTGAGAATG	1200 15	TOCCTGMATO ATTACTOCAA AGGENATATG ANGTAAGAGT TCCCTGTTGC ACATGTACCA
2	GCACCTIGGTG AGAGCTIGCT GTGTGCCAGG CTTTGTGCTG AGTGCTGTTA CATGTATTAG	1260	TOCAMANGG AMACHAMAC GITTIGGATT CITCCCCCCA ITCTCCACAF TGTCCTATCT
	TTCCTTTACT GCTGACCACA TTGTACCCAT TTCACAGAGA AGAGCAGAG AAATTAAGTG	1320	TRAGTOCARG COCTITICAC TOTOLARARA RARARARA TRITITITIC AGCACTGGTS
70	GCTTGCTCAA GGTCATGCAG TTAGTAAGTG GCAGAACAGG GACTTGAACC AAGCCCTCTG	1380 20	TICANARICA ACOTITITAT GOTTARIGGT TIRCCACCAA CTOTICAGAT TICCAGITGA
	CTCTGAAGAC CGCGTCCTGA ATTTCTTCAC TAGAGCTTCC TCATCAGGTT ACCCAGAAGT	1440	OTCITABABA TIGCCARICA TIRICIROCA GCARICACAG AIGRITRAGA GCAGICAARI
,	GGGTCCCATC CACCATCCAG GTGTGCTTGG ATGTTAGTTC TCCACCCTCG AGGTGTACGC	1500 25	CCTCTGAAIT CTTTCCCTAA TAGGCAGCCA TTTGAGAACT GCACTAGCTG ACATCACTAA
3	TOTGAAAAGT TTGGGAGCAC TGCTTTATAA TAAAATGAAA TATATTCTAA AAAAAAAAA		AACATTATCA GCTAAAGCCA AAACCAAATA AAGGCCCAGA CCAACATCCT GGCTCTCAA
	ААЛАЛАЛА	1569	AACCTOTICA AAATCATTAA OTGAAAGGCA GTAAAAGCAG GACTOTIGGAT CATGTICACTG
30		30	CAGCTGACAA TGATTAACAA TAGGAGACAT GCAACCCCCA TTAAGGTTAA AAGTCCGAAA
	(2) INPORMATION FOR SEQ ID NO: 46:		CHARCACAC GCATCTCTTT ATTGGGGAAA AGTGAGACTA TTATGCATTC TTGGTAGGTT
35	(i) SEQUENCE CHARACTERISTICS:	35	TGCAACCTTG CATGAAGAGC ACCCATTGCA TITTCTTTCAT CTITTCAGAAA GCACCGGTAT
			CTGTTCCAAG GGCCTAACAG TACGAAAATA CATTCTGGCA TCACACTCT GAACCCAAGA
9	(C) STRANDEDKESS: double (D) TOPOLOGY: linear	40	CTGTTCTCAT TAAAAATAAT TTTGGTTTGT AACAAAATTA TGAAATAAT TGCAAGCACC
9	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:	2	TOGGIATAGE ATTATTACTG AAACCACTTA ATTCCCAGCT TTTTGAGTTT TTTAAAAAA
	GOCCCCCC MOSMKITHIT TITITITIT ITTAATTAGS ABARTGCCTT TATTAACGAG	09	COCACTOCAC TAAGATTCAC AATTCATTGC TACATACAAA TTAAAGCTAG TAAGAACACA
4 .	AANGAAAGGT TCATTCCTCC TTCCACTCCT TCTCGTTGGT TTTCTGGACA CAGCTCACCT	120 45	CHARGECAC AMETITICECA TECTAAAGEG CAAAAGCCTA AFCATCEGAA AGEGAACAGG
	GNICCIOCIA GAAACOTIGI CAGICIOCITI GIGGOTICCC ICCITGAITG ACTCACOCIG	190	GTAA
Ç	TGTGATGTCT TGAGAAGTAT CTATCCACTT CATGTGAATG AGCACTCCAA TATCAGCCAA	240 50	
3	CATCAATCAT TCTTACCTAA AGAATAATAA GAAAAAGTTA ATATAAAAGA CAAGGGTATA	300	(2) INFORMATION FOR SEQ ID NO: 47;
	ANATAAAGGT TTGAAAATGC TAGTCAAGTT CAAAATTTAA AGAGTAAAAA TCCAGAGATA	360	(1) SEQUENCE CHARACTERISTICS:
55	aagattoogo gtaagttaca ocataaaaa atrogaagaa acttcatgot goggoggaaa	420 55	(A) LENGTH: 475 base pairs (B) TYPE: nucleic acid
	TCTAAAATTA TTCTTACATA AAATAAGTAG ACACCTGAAT TAGAATGAAA ACTGTATTTT	480	
9	CTTTANATG TAAAAGGCTG ACTCTCAGTT TCACCAGTCT GAGCAGAAGT TTGACTGCAA	540 60	(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 47:

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PCT/US98/04493

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8 S 50 3 6 35 မ 25 20 5 5 S GIAGAAGCAG CAGGIGATCT TAACICCITT CAAAAGAGCAG GCCIGICIGG GAAGCCAIGI GCGCACATGA AGACCAAAGC CAGGACCAAG CCCCMASCCT GCTWAACACG GCAGARTCTT GAGGAAAAGG ACTAATICAGA GGAGCCAATG AAGTCACTCC ATGAGTTTCC TGAACCCTGC TOGAGACICA GGTGGGCACA GAGAAGGGTG GAGAGAGAT CTGGGAAGAG AAATGGAGAA TICTAACACA GCTGGTATTT CAAGTCTCCT GGGACCTCAC TCAGGAATGA TACCCCCTCA TAGGTOTCAG CCGCCACCCC CCCCCCATAT GCAGATTTAC TSGGCATGGT AGTGGCCAGC (2) INFORMATION FOR SEQ ID NO: 49: GCCCAGODMA CYTCTGTGAR AATCTGCTTC CCTCCACAGC TGACCC CONSCINGNG ATTANOSTYT GACONTOLAC STASSACACT STOCKGATOS CTACTIOCTS ATOTOTOANA TIOTOMOCIT ACTOTIOTOC CIGOTOGGAC CIGOTOGGA IGAGAGANGG 3 TAAGTAAACT CCCTTTGGAA CTACACAGGT ATGTCTCTCC TTCAACATGT GTGAA TAAAGITATI CIACIAGIIG CAAGITAAGI GIITCIGITI GIICIGCIIT CCIGIIAGCA TAAGCAGCAC AGTGTTATTC ATTTCTGTAA ATTCCTATGT AGAAGGCTCA GTGTTAGAAA AAGGGACAGA GACCTOGATT CAGATCTCAT TITACAATGA AGACCCCAAT GCAGAAAGTC ТОСТОТОВОВ СССАДАААНС ААВВДАССАВ ТВААААСАМС СССАДАДАСТ ТОТАТССВСС INFORMATION FOR SEQ ID NO: 48: (<u>x</u> 5 E Ĕ SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: TGCCAAATTG CTGGTGTCTC TCTTTGGAGA AACCAACCAG ATACATCTGC AACTIGCTIGTIC ATGAAGGAGG GGCCACCTTIG TAAGAGACAT CATTACTACC TITECCAMITYC TGAGCCCTITG AAGGGCAAGG AGGGAAACAG TGTTACCAGA SEQUENCE DESCRIPTION: SEQ ID NO: 49: SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 1366 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 346 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 300 180 120 180 120 475 420 360 8 8 300 240 180 120 25 20 3 8 35 30 2 8 SS 8 5 GOCCTTACTY TITCCTCCCAC AAAGGAGTCG CAGCCACGCY AGCTCTGACT TGCCACTGTG TOOTOGGCCA GGCCCCTGCA TGGGAAGGGA GCCTGCTGCG GGGCAGGCCA GCTGGGGGTG GTCAGCCAGA GTTCTGAAGG CCATGCTTTC AGGTAAAGCA GGATGATGGT GTTTTAAGAC CAGAGCTTGG GACCAGGGCT CCCGCCTTGG GIGAGICIGA CCRYGAGSCG GRCCCCTICA CCITGGCTGG GCTGGTCCTG GTCCTTAGGT TOCTICTOCTIC CCTTCCCTGG CAGTGTTCTG GGGGTGGATT CCCTACAMCT AGATGTTCAA TIGICTICCT GTOCCAGOGG ACCGTGGAGA AAGTGTCAGG GOCCGCTCAC TGCAGCAGCC TITEATIGGECA GEGEGAAGTE GECEAGGTEA GECAGGTGET GECAGEGETE TETETEGGAE AGTOCOTOAA CCTCCCCACC COAATTOCCT CAGTTOTCCT GAGCCTCATG TCTCTCCTGG TITIGGITTAG TCATCIAGAG TCGTCTGGAC TAAAGGTCTT TCAGGTCTCC TTGCCCTGTG GOCCAAGCIT CTATIOTAAC AGTAGGCACA GTATAGICGG ATCATCACAT CAGCIGGGIT CCTCAGCAGG CACAGCAACC CCTCTGGAAA TGGATCACAA ACTCACTTCT CAGCCAGGCA TTIGICAGGT TGICCTIGIT TGGATCCCTC AACTAGGIGA TAAGCACTGG CTCACCTATG CGCAATGANA GITATIGAAG GACTGOTTGT IGATGITIGGT GAGCGTATCC CATTITIATOC TAATAGATIC CATTCTAGGG CCCAGCCGIC TCTTGACTGA TOGTGTICCC GCAGTAATTC CTGTTAGCCA CTGCATCCAC CAAAACTAGT TTATTTTTCC CCTCAAATTC (2) INFORMATION FOR SEQ ID NO: 50: AGCTGTGAAC ATTITICICIC CIOGIAGCIG AACAAAGGIC TAAATIAGCI TAACAAAAGA ACAAAGTICA CGTAGCAGGI CTAGGCAAAG ACTGGGCAAT TGAGCAGAGG AUGATITITA CONCIOTIAC AAAGOGAAIT TIGCIGARAG CICTITIGOGI CCCACIGIIC E SEQUENCE CHARACTERISTICS: AGCAGGGGT TGTGTGTCTG TTCTGTTTCT CTGCTTGCCG ACGIGITITCT TRANCCICAT CCATATAATA GGGCCGIGGG SEQUENCE DESCRIPTION: SEQ ID NO: 50: (B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear (A) LENGTH: 1405 base pairs TTAGGGGGGCC TOGCCACAGA ACACAACCAT AGTITICCCTT GTTGACAATT AACTITICICA CTTAGGCCTG GCTCTCCAGT ACAGGCTGCC CCTACACCTA ATGGTTGTAG AGGGGGATGA AGACOGACCT

> 1020 1080 1140 1200 1260

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720 780 840 900 660

360 420 480 540

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120 180

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	273			· .		
	TITAACCCTT GOCATOTATA ATAGAATITIT GOTGAATGAA AGAACCCAAA TAGGCCAGAT	240	_	TOACCTABOA ACATATOTOT GOOGICTIOTO CTOCTOTOAT AATGAAGACA TAGOOGATTIC	240	9
	AGTCCCCCCA GGCCCTGATA TCCATAAAAG GCTTGGGAAT GCATTATGTA ATTGTCCTTA	300		TOTOCCOGGO CCCCTTGCTG ATGCTCCTCC GGGTCTGCOT CGGGCGTGGG TCTCTGGGGA	300	9
5	CICITITICI TCITITIGAA AAAAAAACA AGATGGGCTC AGATGGATGC CTACGTAAAA	360	'n	CCCTCCAGAG GTGCAGATGS GCTGATGGCC TGGCTGCCTG GTGGTTGATG	360	
	ATGCTTCCTA GCTGTGTACT CATAACTTTT CTTTGAATTG AGTAGTGAAA GGAAGGAGGA	420		CCCTACCTTT TITITIGAG TITATICIGA TIGARITITI TICITIGOTIT CIGGARAAAC	420	
9	GGAAAGGAAA TTAAATGTCC TTCTAGTATT CTCTGGACTC AAGTCTGACA TATGAGATAA	480	01	CACCONCIOS GOACAGEATA ATRABACATG TAATATTITT AAGAAGGAAA AAAAAAAAAAAAAA	480	
2	TAACCTATAT TGAAATGCCA AGAATTGTAT CTGAAACAG AGAACAGTTT GACACATTTA	540		ABABARTINI COCOCOON CASA	3 5	
	TCARGCCTIC ATATTACATA TTAACTGAAA CCAAFTAATA AACATATGAA ATARCCATTG	009	•		Š	•
15	CACAAGGCAA AGCACTAA ACCTITITOTT TCTTTTTCTA CATAGCAGAA ATTGATTTTT	099	15			
	TITITATITE TITAGGGGA CCTAFATAAT TATGACCCAG TGATGTCTIT TGGTGACTTA	720		(2) INFORMATION FOR SEQ ID NO: 52:		
ę	ACCITATICAA ITCACOITAC AAITGAGITG AITCIAGATG GITACIACCI TGAAAAGGAL	780	20	(1) SEQUENCE CHARACTERISTICS:		
3	GTITGGTGCCT TAITGTGACAC GAGCCAGAGC CTGCTGGGGA ATAAACAAAG CAGGTTTCAT	840	2			
	OCCARCACEA ACTEGINGET TTAGTISGICA GATISGICAGT GOTTEACAGA CITECEAAAA	006		(D) TOPOLOGY: linear		
25	TOTGOGGCT TYGGANTITY CCACACCATC CCACOTOTOT TOTTCATACT TCCTCTTTAC	096	25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:		
	ACACTICITIOG ATGGATWAIT TGRAAATOOT GRAAWMMICY YYKRAAITTIG CCCAATAGGC	1020	-	naagtatett goccagitta tiacagaga cgataaatga ticcatgigg ataggocata	9	
ć	WIGROCCACC ATTETTWATG ACACCATAAC CAAATACITC CWIAATOTTG AAATATTAGA	1080	, 6	acatacagag aatgagacta toccagaaat gogaggagoc atttgaaaca acatgagtat	130	
Š	AACCTSTTAC CACCCYKSNA KTWACCOMNA WITTTCCCAT GITTGTGGAA TITGATAITGA	1140	_	CTCAGGGACA GATGGATTGA TTCTGCTATT GGTAGGCCTG GAGGAANGG TCAGAAGTAG	180	
	AATHOCHOSG CTAAGGAATT ACTGSCAAGT TTTAGCCTOT GOSTAATACC TTAGGSTTAT	1200	Ü	CAAAAATGG ATACCAAAAG CACTATTWGT CACCCAAGCT AAGTGGAATA GCTGGCCCAG	240	
35	TIAMATATIT GIMITITIAT TIMMATGTIC ATGAMTGTIT GAMAGGAACA AMATTAICAG	1260	35 1	THGGHGAAAT GCAGGTTTTG CTCTACACTA AGTTCTCCAA CTCTTGATAA GCCTCCAAAA	300	0
	GANGGCTCT TIGCCARGGS TCTTATTITC ACCCTCTITT CIGTAGAA MAGAACAAT	1320	•	acadatista gggaadada acgcagctigg ttatgaadag atatatctica tttcattada	360	•
Ş	CICHTAANGI AHTHHAAG THTHIGGIAF AGTINCIAAF TCCAATTITA AFALAAGITF	1380	, ,	AAATCAATGT CAATGCTGTT AATAGAATCC TTTTATCTTC AGGACAGAGG CAATGCCCTA	420	•
5	twirtaaaa aaaaaaaa aaaa	1405		AICANACACC AGCTCAAGAG CCTCTGATGC CAACCTAGAG GGTACCCAAA CACAAACTTA	480	
			G	GCATAGAGGT AAGAATCTCT ATGTCTTTTG GTGGAGGCAA AGCCATTTGG TTGGTACTTC	540	
45			45 4	ACAGGAACAT CITTCTACCA AGTCTTCATC ATATGGTATG TGCCACGAGT CTCCAGTTGT	- 00	•
-	(2) INFORMATION FOR SEQ ID NO: 51:			ITSCACCACT STSTCATAGE TEAGAATACS CTGAAAGSTT AGTTITGATC CTSGAAACCT	099	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 504 base pairs		•	attracaatt occapotrat stocctocts ccacttaaaa aaggettiggg tctggcatag	720	•
20	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		_	GOAGAMAGOC CTOTGOTCC CTCGTGCCGA TTCTNGGCTC GAGGCCAATT NCCTTAT	7.7	
	(D) TOPOLOGY: linear					
?	(xí) SEQUENCE DESCRIPTION: SEQ ID NO: 51:		55			
3	CGGATTTTCT AGGACCCCAA AAAAAAAA AGGGWAAAA AAACCCKCAA AACCANCCAA	09	_	(2) INFORMATION FOR SEQ ID NO: 53:		-
	aaccccaaaa aaaaaaaaaa tocacaaaaa caaaaaaact aitaaaaaga aagaattaaa	120		(i) SEQUENCE CHARACTERISTICS: (a) LEMOTH: 602 base nairs	-	
99	AACTITICAGA GAATTACTAT TTACTITATT AACTTAGGA TTTATTATAT AAATATATAT	180	09	(B) TYPE: nucleic acid		

	WO 98/39448
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CTTAAAAATA CGAAAATAAA TTTTACAAGG TTCCGGTTTT GGTGGTGGAA AGAGTAAATT CARGOTTIGG GATGAGCAGG TCAATAGITT TGAGAGGGAG TITIGITICCIT TITITITITICT CATTITICGAA TACTGTGGAA GITTTATCIC TIGCATATAC TITATACGGA AGTATTACGC AAAAAAGAAA AATTAACTGT AGCGCTTCAT TATACTATTA TATTATTAIT ATTATTGTGA GCCAGTTTTT TCTTACAGTG ACAGTATCCT TACCTGCCAT TTAATATTAG CCTCGTATTT AACCITGCAT ACGCCTITIC TATCAAGIGC TITAAAATAT AGACTAAATA CACACATCCT CATTATACTC TTAAATTGTT GTCAGTTATC AAACAAACAA ACAGAAAAAT TGTTTGGAAA ATGACTACAG TGTTATACCC TCCAATCTTT GCAGGTGGGC ATGGAACACT GCTTGTATCA GATTATICCC AGGACAGCIT CAGCAAACAC TACAAGICCA CGGIGGGAGI GGATITIGCI CCCGACCACA CTTCCCGCCT CCCTAAAACG CACACCCCGC TAGCCATGGG CAGCCGCGAC CUTCACITICA GITTIGAAGAG GGTCCGGATC CAAAGGGGTT AAAACGAGCG AACCCCGATC AGTCACTGAC TIGGAGCCGC ICGGGGGAAG ICCCGCCCAG ACAGGCGGIG GGIGGGAAIG TICTCACGIA TAITIACCIG IGACIIGIAT TIGITAITTA AACAGGAAAA AAAACATICA ATTATOTTO ACOTTACCAA TOCCACTACC TTCAGCAACA OCCAGAGGTG GAAACAGGAC CTGAAGGTTC TCCAGTGGTC TGACTACGAG ATAGTGCGGC TTCAGCTGTG GGATATTGCA CACCIGITCA AAGIGCIOGI GGIGGGGAC GCCGCAGIGG GCAAGACGIC GCIGGIGCAG GOCCAGGAGE GETTEACETE TATGACACGA TIGITATIATE GOGATGEETE TGEETGTGTT CTAGACAGCA AGCTCACACT ACCCAATGGA GAGCCGGTGC CCTGCCTGCT CTTGGCCAAC AAGTOTGATO TGTOCCOTTG GGCAGTGAGO CGGGACCAGA TTGACCGGTT CAGTAAAGAG 600 540 480 420 360 300 180 180 240 120 600 480 420 360 300 60 6 330 23 8 2 5 55 50 25 8 CCCACCIGAC CATTITATIA AGIACATIIG AATIGICICC IGACIACIGI CCAGIAAGGA GENTITITICA AACITICIAAC GICATAATTA AGETTICICIT GICTIGGCAT CAAGAATAGI TICICATCAG CCCICAATIT GIGAICCGGA ATTITIGIGAG AAGGATIAGA AATCAGCACC GOGCCCATTO TCACTINGAA ANGACACCTO GAACCCATOT GCATTTCTGC ATCTCCTGGA ACCCAAGGG ACTACATCAA TCTACAAACC AAGTCCTCCA GCTGGTCCTG CTGCTAGTAG CGAGATCATG CCACCGCACT TCAGCCTGGG TGACAGAGAA GGACTCCGTC TCAAAAAAAA GGCAGGCOGA TCACATGAGG CCAGGAATTC GAGACCAACC TGGTCAGCAT GGCAAAACCC CAAGITITIT GOCCGOOCAT GOTGGCTCAT GCCKGTAATC CCAGCACTTG GGGAGGCCAA TTAGCCTTTC ACATOTISCT GROTCACATT AGTGCCAGTT AGTGCCTTCG GTGTAAGATC IGITIGGCIT ATTTICCAIC CCAGITCIGG GAGGICTITT AAGTCICITC CCITIGGIIG ATGAGAGICC TCATTGAAAA GATGATGAGA AATTCCACAG AAGATATCAT GICTTIGTCC AACGGTTTCA CAGGTTOGAC AGAAACATCA GTCAAGGAGA ACAAAAATAT TAATGAGGCT GOGTOGCCTG TOGTATIATOG AAAAGTAGCA GGGTGGTCAG GGTGGGAGAC ACAAGATGTT CCICCIGICI ACIGGCICCA AATAGACCAI GICAGCIICA CCCCCIGGCI IIGIGIGICIAI CATTARATTT CTTACAGIGA ACTACATATT GICCATAAGT GCTTCATCAG GACTCATCGC AAAGAGATGG GCTCTTTATT TICTCGAAAA ACCAATTIGG AGTTACTCAT TITTTCCATAA (2) INFORMATION FOR SEQ ID NO: ACAATCNAA AAAAAAAAA AAAACTCGAG GGGGGCCCG GTACCCAAAT CGCCSTGATA GTGATCGTAW TACTOTOGAG ACTUAGOTOG GAGAATOGOT TGAGACTOGG AGGCAGAGOT TGCAGTGAAC COTCTCTACT AAAAGTACAA AAATTAGCCA GOOGTGATGG CACGTGTCTG TAATCCCAGC AGAAGGAGGG GTCTGGGCAT CTGTGGATTT TTGGCTACTA GAAGTGTCCC AGAAGTCACT IGIAGGCCGT IGITICAGAT ICTTICIGIC TIGGAAIGTA AACATCIGAT ICIGGAAIGC TIGATATCAT GACTICCAAT IGAGAGGAAA AIGAGATCAA AIGICATITC CCAAATITICI IGCGTTTTAG AGATCATAAT TCTCACCTAC TTCTGAGCTT ATTTTTCCAT TTGATATTCA E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear (B) TYPE: nucleic acid
(C) STRANDEDNESS: double ٤ LENGTH: 1896 base pairs 55 1740 1320 1680 1620 1560 1500 1440 1380 1260 1140 1200 1080 1020 960 900 840 780 660 240 180 120

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SEQUENCE DESCRIPTION: SEQ ID NO: 54

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INFORMATION FOR SEQ ID NO: 54:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1749 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 53

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TOPOLOGY: linear STRANDEDNESS: double

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TITIATAGECT AGAGCCTITIA AAAAACCCAG CAGAATGEAA TECAGEATET GITTATTIGGC

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AGAGITICAG		1896	ANCOCCOGGG NGGGCCAGNA GGTTTGCCAG TTGAGT
GTGAAGATGC	55	1860	GIOCCIGITA AICTICAGCI ACTINGGGA GGGCITGAAG CCAGGGAGGA ACTGCCCTGG
TIONION		1800	GAAACCCCGT CITCITACIT AAAATACCC AAAATTAGGC CAGGGGCGTG GATGGGTGGG
GATTGTGATA	50	1740	ACACTITAGA GOTTIATIAT ITITAAAAA CITITCITGA ACICCIGGGC CAACAIGGGT
TAGAAATTAA	•	1680	actiggataat gtggagtgac tagagaagtc ayatatcact gtaagataca gttaggagta
GATGGTGCTG		1620	TGCTTCTAGA AGTCTGCAAT TCTTTAGTTT TCTTTGGTGC ATTATTATCT AGGTGCCATC
TGTTTTAACC	45	1560	ATCCATTTGC ATAATGACCC AGATCATCAT TITCTGCAAC TGAGAATTAT ATTICATCAT
TGCAGATTAC		1500	CTTCCATGCT ATCGGATGTG TTGGGCTCCA TGCAAGAACT TGGAAGAAA ACAGGCAGGA
TIGGICIAIT		9	TSITATTAGT TSITITSTIT ATARITITAT AAAATTAIT CALGOSAGI TIAASITUCA
AAAGTATTTA	ć	3	AMMGAMAIA INCICCIMA CITTACIAC CACILCIMA ITMATOTOS ALLICAIMA
GGAAAGGGTG		1320	ogatangana tcacaggnga tagcagtigt cattcagtaa tittcctagna ocagcagooo
GTTTCCCATT	35	1260	TICITICICIA IGCCAIGACT IGAAAAAGIT IGGGAAGCIC TITAGCAATA ICAGCIAAAA
TATATGITIA		1200	TOGGGATCCA GCTCAGAGCA ATCTCTTGCA TTTTTTTACC COTGTATGTA CAGATATCAT
TATCTACTGA	30	1140	TGACTOTTTA GTTCACAGGT TCCCATTGAT TGTGAGCAAG ATATTTATCT CTTTAGCCCT
		1080	AAGGAATTTA CAGTGATTTT AGTGCTTGTC AGCATTTTTC CATGAGGACT TTCATACATT
ATAGAAAGTA		1020	TATTGACATO TAAAAGACCA AAGTAATTTT TCTGAACTTC TGCAATTCTG AGAACTCCC
ATCARCAGE	36	960	TTGACCAGGG TCTCACACCC TGGAGGAATG TTAAGTAAGA GAAAGAACCT CTTTCCTGAA
		900	GOTCAACTIG ACCCIGCCAT GITGGITTGA CITACTAAGA CACAGGAATC ATTGTTTTCC
CTABABTACC	20	840	CCTACGAGGT TAAAGAACAC ACTGTTCCAC TGTATGGCTT TGGCCCTGAG TGGCCAGGGA
THECTTARA		780	CTIPAAGAAG GGCAGCCAAG GIAGIAACCT AAAAAIAGIG CCCAGGCATA IGAGAGIIGI
Telephone		720	AATITICCAA CIAGAGGAAG AGAAACTIGI GGAAAAGTIC ITITITITIC ITITITITI
Thirting	<u>×</u>	099	GTTTACAAGT GTTTGAATTT GTGATCTGAA ATAACACAAA ATTAAAAACA TGATTTCTCT
TCITITIANA		909	CITITIBACC GIMARINICA GANIGAMICC ICTICCCAGG GANITGAACA GAAGCTIMAT
(TX)	10	540	ATMATACTAG TAGGAAGTTA ATMACTGOTT CTCTGTGTTC CAAGCACAAT ATTACAACTT
3		480	TTOTTTCCTT GATCTOTCAG AGGTCACAGT AACCTGGGCC GAGCTGTTAT TATTTATTAT
		420	OTTINITIANA AGGANAMAN TOTOTTIOCHA TOTOCTOTIAN TCACARGAGO AGANAATHAC
3		360	TETTITITIES CAGATIVETIS AAATTAAANG AATTGAAAGG GAAACTCAGA GTACTAGGA

(2) INFORMATION FOR SEQ ID NO: 56:

) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1753 base pairs
(B) TYPE: nucleic acid
(C) STRANDERNESS: double
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 56:

9 120 180 240 300 360 420 480 540 9 99 720

ATCOCTCACT GATACTATCT ATTITCTTIAG TAAGAATGTG TTAAAATTAC AAIGAICITI TAAAAAGAIG AIGCAGIICI GIAITITATIG IGCIGIGICI GGICCIAAGI ATAGACATTT GTGGGGCTCA CACAATATAT GAAATAGTAC CCTCTAAAAA GAAAAAATTT AATATCAAGG ACTATTACAT ACTTCATTAC TAGGAAGTTC TIACTITICIT ICATATIGGIT TITIGGITCAC TGGCTTAAGA GGTITCTICAG TAACAGAATC TGGOTCACAT GAAAAAAAT CATTTTAICC GTCTTTTAAG AACTETITET GACCAATAGT GETGGCACCG TIGETICCTE TITGGGAAGA TOTCAACATG GCTAACAATC TTCAAATACC CAAATTGTGA TAGCATAAAT HITTAIGCCT CAGIAININA TIAITTAAHT HITTAGGIAA IGCCTAICTC ATGITITIAC AGIGGCCTGC TAITIGAGGAA AGGTATICIT CYATACAACT TITGAGAACA TIGACAGAAA TIAIGCAATG GITIGITGAG AIACGGACTI TITAATCAGT TIGCTITCCAA AGTGGCCTAC TCAAGAGGCC CTAAGACTGG AAGGAITICA AAAACITICI AITICCITICI TAAACCIACC AGCAAACIAG TITICAATITIG AAATCATICT TAITICCCTIT AAGAATGITT AIGTAIGAGT TAGGGAACCT ATGCTCAGAT ATTCATGGTA AGTCTCCCTT CACCTGTTAC AAAATCAGGC GGTCAAACTT AGAGCAACAT TGTCTTATTA AAGCATAGTT GACACTTAAA ACAATCACTG AAAACTTGAT CCACATCACA CCCTGTTTAT CATCTIGGAA GCCIAAGCIT CIGAGAAICA 1G1GGCAAGT GIGAIGGGCA AGAGAAGATG TITAGTAGCA ATTAAAGGCT GITTGCACCT TTAAGGACCA AOTGATTCCT GOGGCAGAG TGGCATTATG TTTTTACAAA ATAATGACAT TITICCATOTT TOTTTGCTTG TTGAATTTTT GAACAGCCAG TTGACCAATC GCCACAGCAG CATACCAGTT TCCATCCTAA TAGGAATGAA ATTAATTTTG AAATAATAAT ITATGTGTCT GCATATTGCA GAACAGCTCT GAGAGCAACA AAGGAAAGAA GCAATCAGTA GAGAATTCAG GATAGTTTTG TITTAAATTCT

780 840 900 960 020 080 140 200 260 1320 1380 1440 200

	WO 98/39448
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1020	960	900	840	780	720	660	600	540	480	420	360	300	240	180	120	60							1753	1740	1680	1620
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		ARARAMANA ARANCTOGRG GGGGGGCCC	ACAAGTGGTA GTGGCATTCT ATTTATTGTG ACCTTTTCAA TAAATAGATT TAAGTAAAAA	CTTTTTAGC ACTOTTTTG TTTTAATGGT ATAITTTAAT TGGCTACTTT ATTGTTTAGG	GACGTCTCCT CTGCCTAGGG AGCAGGACTT GGGCTTAGGG CAGGTGGAAA AAATTCCAGA	TTATOCCGAG AAGATCTCAG CTOGATOCCA ACATOTTCCG ATGCCTGTGG AAGACATGCC	CATTHAGTOG CACAMARICA GAGCAAGAAA GOGATOCCCT TCCCAARTCT CTCAARCCTT	TICCCAAAGT GTOGGTOGGT CCGTTGGTTC CCGAGATACT TITHGGTOGT ATOGGGCCTG	CAGGIGIOIG TITCCCCTIT GIGITAAGCG IGAGGCAGAG GGAGACGITA GICCCAGCAT	ACCACCCATA TITICOACCC ATCICCTCCTC CCCTTCAACC TATGATCAGA GAGGGGACCA	ACCICIGAGG GETGATAGGG GIGGGTTIGT TGAGAGGGAC TIOCIOGGCC TIOGIGIGAG	TOTTOGRAD CICCICCIOS TIGOCCITOS CIACCIGAIS TCCCACCIGA GICAGCOSIS	CAACCATOCT GTOGAGCCOG TGACCTCCAT CCTGCTCCTC TTCCTGCTCA TGATGCTTGG	COCCUTICAAT GACCICAACC GGCAGCIGGT GAACAIGGGC ITITCCGCAGI GGCATCICGG	COCHAGGOST COTAGOCCOC GCCAGCAGCA GGCAGGTGCC AGGCTGGGTG CTGCTCAGTC	CCAGCCACCA AGAGGTAGCA TICCTCGACA GAGCTICTIC AATAGGGGCC ATGGTGCTCC	осодитство оссилложно импестоси осиссистве ссессийтся осиссистис	CAAGOCTOCC TCAATTTCTG GTGCAGCCCA TCAGGGACCC ACAGCGCCTG GGAGGATGGT	торорство адасьська тепнетенке второтенсе тетрговыем телетомето	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1049 base pairs (B) TYPE: nucleic acid	(2) INFORMATION FOR SEQ ID NO: 58:			TCAWAAAAAA AAAAAAAAAA
		1049	1020	960	900	840	780	720	660	600	540	480	420	360	300	240	180	120	60							1220

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CATOTOATTA CITCACICCI GGACIGIGAC ITTCAGIGG AGAIGGAAGI TITTCAGAGA

ACTGAACTOT GGAAAAATGA CCTTTCCTTA ACTTGAAGCT ACTTTTAAAA TTTGAGGGTC

TTACOTOTOC ACAGAGAGOT CACCTTTTTC AGGACATTGC ATTITICAGGC TTGTOGTGAT AAATAAGATC GACCAATGCA AGTGTTCATA ATGACTTTCC AATTIGGCCCT GATGTTCTIAG

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TOGACCAAAA GAAGAGGAAT ATCAGGTGA AGTCAAGATG ACAGATAAGG TGAGAGTAAT
GACTAACTCC AAAGATGGCT TCACTGAAGA AAAGGCATTT TAAGATTTTT TAAAAANTCTT
GTCAGAAGAT CCCAGAAAAG TTCTAATTTT CATTAGCAAT TAATAAAGCT ATACATGCAG
AAATGAATAC AACAGAACAC TGCTCTTTTT GATTTTATTT GTACTTTTTG GCCTGGGATA

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(i) SEQUENCE CHARACTERISTICS

(2) INFORMATION FOR SEQ ID NO: 59:

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SEQUENCE DESCRIPTION: SEQ ID NO: 57:

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(2) INFORMATION FOR SEQ ID NO: 57:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1220 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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AAAAAAAAAAGG GGG

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TOCTTGAGAT TICACTACCT TIATGITAAA AGTIGIGIAT AATIGITAAA ATCTGTGAAA

GUAGCCARIT ARACHAGITI CATRIGIRIT TITICCAGIGI TURATCICAC ACACIGIRCI TURADAGATI COTTICCATOC TURATRACAR ATRABAGAGO CORTATRIRI TUCCTOCTIR

1560

TUGGTTTTAA ATUGACATTU TCTOTACCAG CTTCATTAAA ATAAACAATA TTTOTAAAAA

1200

СВАТЛАВАЛАС ТССАТТТАВА ТТЯВЛАВАЛА ВАЛВВАЛАВА ВАВВАЛАВАВ ВВАЛВВАВАВ

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≱	WO 98/39448 PP	·CT/US98/04493	Š	WU 98/39440		3
	107					
	(A) LENGTH: 1776 base pairs (B) TYPE: nucleic acid		8	GGGTHINING ATTOTATIAN AANAANAAG GTHINININGC NININICINI AINTANINIG	1680	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear			acocagaat aaatctatga gaaatctatc tacaaahnaa aaaaaaaaa aaaaaaaaa	1740	
S	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:		٠ ۲	AGGARITCGA INTCAAGCIT AICGAIACCG ICIAGC	1776	
	AAAGAGGATG TGWAGCTAGA GGTCCCCGAT GCCTGGTCGG ATGGGAAGCA CAAGGCTGAG	. 09				
01	GOACTGGATT GTAAAGGCAC TAAGTCGTTC TGCGGTGAGA ATCAGACATG GGGGACCTCT	120	2	(2) INFORMATION FOR SEO ID NO: 60:		
	ACCITICACAT CCTCITITICCT TOCAGSTICTG CACATOCTICA GCCCAAGTOC CCCACACTCA	180	-	(1) SEQUENCE CHARACTERISTICS:		
ÿ	GTOCAGTGAT GAGTGCGGAA GTGAAGGTGA CAGGGCAGAA CCAGGAGCAA TTTCTGCTCC	240	15			
2	TAGCCANGTC GGCCAAGGG GCAGCGCTGG CCACACTCAT CCATCAGGTG CTGGAGGCCC	300		(C) STRANDEINESS: double (D) TOPOLOGY: linear		
	CHOSTIGICTA COTOTITIGOA GAACTICCIGO ACATIGOCCAA TOTTIAGAGAG CTGOCTIGAGA	360		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:		
20	OTGACTITIGG CICTACCTIC COGCIOCICA CAGIOTITIGG TTAITGGACA TACGCIGACT	420	, 20 8	acagataaat aaataaataa taaattaaat taaataaaaa atctgagcta atctgaataa	. 09	
	ACTIAGCIGA AGCOCOGAAT CITOCICCAC TAACAGAGGC TCAGAAGAAT AAGCITCGAC	480	٠ ٦	ATTENCIALAT TTCACATGAA AGCCAGGATT TCTGGCTTCC CAGGAACAGT CAGAAGAGCT	,120	
25	ACCICICAGI TGTCACCCTG GCTGCTAAAG TAAAGTGTAF CCCATATGCA GTGTTGCTGG	540	25 AG	AGCINGCANC ACTIGATICIGC TIGGCTACCT TOTITIGGAAC AACATGAAAT CTAGOTICCT	180	
ì	AGCCICTICC CCTOCGTAAT GTGCGGCAGC TGGAAGACCT TGTGATTGAG GCTGTGTATG	009	Ę	THITHTHE THITHGGCCC ACTICATCA TECACATGAC CTGCCTGGCC TCTGCAGGTA	240	
	CTGACGTGCT TCGTGGCTCC CTGGACCAGC GCAACCAGCG GCTCGAGGTT GACTACAGCA	099		AGIGACIATO CAACAAAAT GTAGCACAGG TTTTGTCGCT GAACTACGTG GTTTCAGGTC	0 	
30	TOGGGCGGGA CATCCAGCGC CAGGACCTCA GTGCCATTGC CCGAACCCTK AANAAAAACC	720 3	ი გ	CAGCTOTICC ACTIOCINGC ATGACCTOOF GCCGAATTCC NCCACGAAGT TITTITITT	360	
	ATTANAGITA CGACGGCAGC AGCAGGGGA GCCACATCTC AGGACCTGA GCAACACTG	780	Ē	TITITICAGIS CICCAGICCC CCIAITGGAG AATCCIGCCC CCCCIGGGA CAGAAIGIIC	420	
7	ACTERACTICA GGGAACCAGC TECTIGGCACC AACCAGGGCC ASCCAGCAAG AAAGCCTCAA	840	35 AC	ACCITIGACIC CACCANTICC TGA	443	
C C	AGGGCAAGGG GCTCCGAGGG ANCGCCAAGA TTTGGTCCAA GTCGAATTGA AAGRACTGTC	006				
	GITTECTICCE TOGGGATIOTO GEOTECCAGE TECCTOCCTG CETETIAGA GTECTEAGAG	096				
40	AGCETTOTION GCCCCTIGGC AGCTGATAAT CCTAGGTTCA TGACCCTTCA CCTCCCCTAA	1020	40 (2	(2) INFORMATION FOR SEQ ID NO: 61:		
	OCCCAALCAT AGNICACACC TICICIAGGG AGGAGKCAAA IGIAGGICAT GITITIGIIG	1080		(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2888 base pairs		•
¥	GTACTITICIG TITTITIGIGA CITICATIGIGI TECATIGCIC EXCRETIGEA TACTECICE	1140	45			
7	CITGITICCI TAAGAGCICA GCATCIOICC CIGITCAITA CAIGICAITG AGINGGIGGG	1200		(D) TOPOLOGY: linear		
	TRACCCTIGAT GGGGGTCGCT CTGTCTGGAG CATAACCCAC AGGCGTTTTT TCTGCCACCC	1260		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:		
20	CATCOCTICA TECCTIGATIC CCASTITICTA TACCCTACIC CTGACCTATT GAGGAGCTIC	1320 5	50 TT	ttaatsttot caataaccac cageccaarc agaatttata tgacctggat gaagatgatg	09	
	TRANSPOCCA TAGGGCCCC ACCTITACTC ACACCCTGAG AATTCTGGGA GCCAGTCTGC	1380	¥	ATGGTATAGC TTCCGTTCCT ACTAAACAGA TGAAGTTTGC AGCCTCAGGC GNCTTTCTCC	120	
;	CATGCCAGGA GTCACTGGAC ATGITCATCC TAGAATCCTG TCACACTACA GTCATTTCTT	1440	X X	ACCACATIGGE TGGGCTAAGC AGTYCCAAGC TTTCCATGTC CAAGGCCCTC CCTCTCACCA	180	
ç	TTCTCTCTCTC TGGCCCTTGG GTCCTGGGA TGCTGCTGCT TCAACCCCAG AGCCTAAGAA	1500		AAGTOGTICA GAATGATGCA TACACAGCTC CTGCTCTCCC TTCCTCTATT CGAACAAAA	240	
	TOGCHOCCGT TTCTTARCAT GITGAGAGAT GATTCTTTCT TGGCCCTGGC CATCTCGGGA	1560	ت ک	CCTTGACCAA CATGTCCCGG ACACTGGTGA ACAAGGAAGA ACCCCCAAA GAGCTGCCAG	300	
9	1 ACCITICATIGG CAATCCTIGGA AGGGITTAAT CTCCITITIGT GAGTITTGGTG GGGAAGGGAA	1620 6	ີ ຍ	CTGCTGAGCC TGTTCTCAGC CCATTGGAAG GCACCAAART GACTGTGAAT AATCTGCACC	360	

8 ႘ ဗ 25 8 15 8 45 5 8 S CAGAAGGCAG TAGGACCTGG TITTITCAGGT ACTGGGAAGCC GGGGGCTCAC TGCTTGCACT CCTCCTCCAC TICCCATOCC TCTATOTTAC CCATCTOTOT CTCCTGTGCA GAAGGAGAGG CAGTICTITA AAGAATGCIG CITTITATIC ICCTAACCCI TICAAGTGGG IGCAGACTIC GOCTICTIGAGG CTOCCCAGGA CGGGAAAGTIC CAAGGAAGGG GCCTGGTIGGT GCTCCACTTG TOCAGOGGIC AGICIGICAG OCAGGAAGGA COCAGGAITT GAACCCAGCI TOAGIGIGCA CACTAGIATA ATTIATAATT ATAACCTATT CIGATITICIT TICAAATATI AGGIGICCTA AAAGTGAG CACTOTATIAA GCAGTOATOT TOOGAGACOG GGAGGAGAAG GTOGTOGGOT AGTOCTGTGT AAAGCAGGTA TCTTCTGGTT GTCACAGAGT TTCATTGAGT CCAGCTGCAG CCACGTGGCC TOGITAGOAG CIGGAAGACA TICCICCOAC ACITITICCCI TOCIOGCCCA AGAGAGCATO CTIGAGGGCAC TCGCTTTGCT CCTGTCAGTA CACACTCCCA AACAGTTAAA CCCAGCTCTA GICTACACGI ICAIGACACA TIICCIICIA AAGGIICAAA GICAAGIGII TICIGAAGCA TOCCAGACTO AGTOTOCTAT GTCAAAAAAAC TOCATOAAGO TTTTOTOTGA AGATOCTOTT GITIGOCTIATIG AAGGITTIGGO ACTICATICITI GCACTIGITICO CCAAACTITIG GACTIGAGGIA (2) INFORMATION FOR SEQ ID NO: 62: TGTTATTCCT AAAAAAAAAA AAAAAAAACT CGAGGGGGGG CCCGGWACCC AWATCGCCSK AAGGGGCATT AAGAGATGAA GGGTGATTAT GTATTACTTA TCCATTTCTG AATAAACATT ATCTRIAGET GETRETATAG GTRACCATET GETACATERA GREGACCTET TECCETECTE GIOCTIAGOG TAGOGATOGI AAATATECIC CCIGCATOGC TITATECTICC CICICATECC CAGTCTGATT TCTCCAACCG AGTTGAAATT TCCAAAGCAA GTGCTTCTTT AAATGGGGAC AACTOTOCCA ATTTGATCAG CACTCTTATT ACAAACTTGA TAAGCCAGTA TCAGAACCTA TICGCAGAAT ATATTAAAIG TATCCTAAIG GATGAAAGAA CIITTITAAA CAACAACATI CTCCAAGAGC AAGAAGCCAA AGAAAGAAAA ACTAAAGATG ATGAAGGAGC AACTCCCATT ATTOCAACTO TOCAAGAGOT TITAAGOAAA TOCAAGACTT GTOTOCAACA GAGAAACTOA Œ. (i) SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENOTH: 1851 base pairs 62 2580 .2340 2880 2820 2760 2700 2640 2520 2460 2400 2280 2220 240 180 120 600 540 420 360 300 480

20 5 55 80 S CATCHATGAA AAAGGAGAGC GAGCTGCCTC GCAGGGTGAA CTCTGCCTCC TCCTCCAACC CIGGGIGICT TIGCIGGGG GGGGCIGIGT TOTGAGCCCT CCCGGTICTC ACCICGCCIG GODATIGGADA COCCTIGGTOT COTGACGGGA GOCAGGTICGG COTGAGAGGT GTGCCGCTCC OGGIOGETTI CTIOTOGCCC CATOGGANGC AGCGIOGOGG CTGTCTGAAG GACCCTGCTT CCTCCCCGAG TOGATICAGAC COCCUTTICT GIGITIOTOTT CIGCOCTOTG CICTICICIC TACGITIAACG CCATCGGACT GGAGACCCCT GATTOTOGGA AGGGTTOCCA GGGATAAAGA GCTTCCTCAC CTRIGACCAC GCAGCCCACA GAATICAAAA TCAAGCTITG AGCAGGGGAG TGAGGCAGCC CCCCTGCYGA AGTGGACCCT GACACCATCC TGAAGGCACT CTTCAAGTCC TCAGGGGCCT ACATGAATGG GAATGITATC ACCTCAGACC AGCCCATCCT GCTGCGGCTG AGTGACAGCC COCCATATAA GAAGIACAAC AACCGGISTC TOGACGGGCA GCCGATGAAG TOCAACCTTC CICCACIOGI CCAICCIGGG GIAGCGGAGG IGGIGITIGI GAAAAAGGAC GAIGCCAICA CICCAGICAC TGAGGAGGAC ATTOTIGAGC TITTICIGIOT GIGIGGGGCC CICAAGCGAG CCAGTOCCTC CAGGCCTCAC TAGTGGCAAG GGCAGGATGA GGCTGCACCG CTGGGAAGAG TOTOTOTOTO CHOTOGRAGOT GOOTIGGOTIGG GGAGCAGGTO TOAGGOOTOT TOTOCOTOTO TITICCIOTAG TAIGITTICIT CATCICATCG CCAAGOTAGG CTIGIGITIT TCAGIGIGIG AGAAGTOGGG GCAGAGGAGG GTGGCTCTGT TTCCCCAAGG CAAAGCTTAT GACCAATOGG CCCCATCCAT CCATGACCAG AGGATTATTT TCCTGCCTTG GCAGAGGAGG AGGAGTCAAG GITCACCITY TOTATOTAAG YTOTTOOCTT GGAGTOOOGT GTOGTOTOOR COCAGAGGAA GTTOTOCAGA CTOGAGOCAT ATATCCAGCT GCCACCAAGG GGCACTGTTT GTTCCCACTT ATGTGAGTGA AGGCTCAGCC TCCCATTGTG CAGTGCTTGG GTTTGGAGCT TATTTGAATG GAAGAGGTCA GCACTTAACC ACACCCIGGT TITIOTOTAGC CGCCAGCICT CITICIGGITG GGCCTTIGAA GGTTCTGGGA GCTCTGCAAA ATCAGTAGCA AGTGCTGGAA AAGGCACATG CCGAAGATAC CCTOCTOGIT GOGAGTGAAG AGAATCCAGG CTGGCAGAGC TGGAGCCAGT TGGGGAGCAC GAAACTICAG AGCCCAGGCA GICCCIGAAT GACCAGGCCA GIGTIGICAC IGAGIGGICC BEAGCAGGGC AGCTCTACCA GGCAAGGTGT TTCCCCAGCA TAGGCGCAGA CAGTTGGGAC CCCTTTTCCT TGAGTTOTOC TGAATGCCCC ACCCCAGCTC TCTTTCCCTT CCGAGGGGCT GCCTTTCCTT TGTGTGTATT AAGCTTTTCA AACAATGGAG CCICAGCCCC AAGCIGATIT CITATCIGGA AATGGTACAC IGAATTCICT GOCTICTICCAT TICTOGCCTC AGTIGICIAC AGGACAGIGG TCAGGGAIGC 1260 1200 1140 1080 1020 1440 1380 1320 1800 1620 1560 1500 1980 1920 1860 1680 960 2040

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TCAAGAGCTC CCAAGATTTG CTTGAGGCTA GCCCAGTGAA RAAAACCAGA GACTCATGTT

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	AAAAGCCCC GTCTTAGCAG TGATGAGGAG CACACTGTAG ACAGCTGCAT CAGTGACATG	099
	AAAACAGAAA CCAGGGAGGT CCTGACCCCA ACGAGCACTT CTGACAATGA GACCAGAGAC	720
2	TECTCAATTA TIGATECAGG AACTGAGCAA GATETTECTT CECCTGAAAA TAGTTETGTT	780
	AAAGAATACC GAATGGAAGT TCCATCTTCG TTTTCAGAAG ACATGTCAAA TATCAGGTCA	840
9	CACCATGCAG AAGAACAGTC CAACAATGGT AGATATGACG ATTGTAAAGA ATTTAAAGAC	006
2	CTCCACTGTT CCAAGGATTC TACCCTAGCC GAGGAAGAAT CTGAGTTCCC TTCTACTTCT	096
	ATCTCTGCAG TTCTGTCTGA CTTABCTGAC TTGAGAAGCT GTGATGGCCA AGCTTTGCCC	1020
15	TOCCAGGACE CTGAGGTTGC TITATICTCTC AGTTGTGGCC ATTCCAGAGG ACTCTTTAGT	1080
	CHTATGCAGC AACATGACAT TITAGATACC CTGTGTAGGA CCATTGAATC TACAATCCAT	1140
Ş	STOSTCACAA GEATATCTGG CAAAGAAAC CAAGCTGCTT CTTGACATTA GOTGTAGCAT	1200
₹	STOTACTITIT ANGROCCICA CCCCCAACCC CCATGCTGTT TGTATANGTT TTGCTTATTT	1260
	GITTITISTICS TICAGITIST CCAGISCICT CICCTICAAT GCCAAGATAG AITTATAGGC	1320
22	TIVATICITIS STCAGGOAGA ACTECAGATS ANAMANCIT GCATETICAS TATACTICET	1380
	AAAGGCCAAT CAGATAATGG ATATGTTTTA TGTAATTAAG AGTYCACTTT AGTGGCTTTC	1440
ć	ATTIBATATS GCTOTCTGGG ANGARCAGGG TTGCCTAGGC CTGTACAATG TAATTTAAAC	1500
₹	TRACAGCATT TITACTISTIC ATGATATISCT GICCTCTISTS CCAGTITITGT ACCTTATAGA	1560
	GSCAGATTGC CTCCGATCGC TOTGSTTCTT ATTAICAAA TTAAGTTAC TTGTATACGG	1620
35	AACAACACA AGAAATTIGA TICTIOIAAAG AATECTICITT AGCIGTGGCC TGGCAGTAIA	1680
	TANAISGIGC TITAITINAC AGANIACCIG IGGAGGAANI ANAGCACACT IGNIOTAANA	1740
ç	ATARITICITY TAITITITAIT GACATGACTG ATTGATTGCT ATTCTGTGCA CTTAALTAAA	1800
€	ctgattgtga tgacttmmaa aaaaaaaaaa aaaaaaaaa aaaaaaaaa a	1851
45	(2) INFORMATION FOR SEQ ID NO: 63:	
50	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 3542 bese pairs (B) TYPE: nucleic acid (C) STRANDERNESS: double (D) TOPOLOGY: linear	
ď	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:	
ર	TCCAATGCTG AIGAGCGTCT TCGCTGGCAG GCCAGCTCCT TGCCTGCTGA TGACCTTTGC	09
	ACAGADADTG CCATCATGCT GADACGATTC DATAGGTATC CGCTGATCAT TGACCCTTCT	120
9	GGACAGGCCA, CAGAATTCAT TATGAATGAA TATAAGGWTC GTAAGATCAC ACGGACCAGC	.180

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480 540 909 9 720 780 840 900 960 070 0801 140 200 260 1320 1380 440 200 1560 620 0891 1740 800 0981 1920 980 CTGGTCCAGG ATGTGGAAAG CTACGATCCA GTTTTGAACC CGGTGCTGAAGTG CGCCGAACAG GGGGAGAGT GCTGATCACT CTCGGGGACC AGGACATAGA CCTGTCGCCA TOSTITISTICA TETTECTISTE CACCOGGAT CCAACTISTEG ASTITECCACC AGAICTETST TCCCGGGTTA CTTTTGTAAA CTTCACAGTT ACCCGTAGCA GTTTACAAAG CCAGTGTCTA AATGAAGTAC ITAAAGCAGA AAGACCIGAT GIGGACGAGA AACGAICTGA ICTICITAAA CITCAAGGG AAITICAGCI CCGITTGCGI CAGCIGGAAA AAICICTACT ACAAGCICTG AACGAGGTGA AAGGGCGCAT TTTGGATGAC GACACGATCA TAACCACTCT GGAGAACCTG AAGAGAGAG CTGCAGAAGT CACCAGGAAA GTTGAGGAGA CGGACATTGT CATGCAGGAG CAGCGCCTGT CCATTATAAC AAAGGACCTC TTCCAGGTGG COTTTAACCG AGTGGCTCGA GTGGAGACCG TGTCCCAGCA GTACCTCCCC CTCTCCACCG CCTGCAGCAG CATCTACTTC ACCATGGAGT CCCTCAAGCA GATACACTTC TTGTACCAGT ACTCCCTCCA GTTTTTCCTG GACATITIATO ACAACGICOT ATACGAGAAC COGAACOTGA AGGITOTOAC CGACCACACA GOCATICCTIC ATCAGGACCA CAPTACCTTY GCCATGCTICC TGGCAAGAAT CAAACTGAAG GGCACCGTGG GGGAGCCCAC CTACGATGCA GAATTCCAGC ACTTCTTGAG AGGAAATGAG ATTIGECTION GEOCTOCOCE CACCECCAGO ATCCAGOGOC TGACTIGEGA GCAGGOGGAG GOGGAGICTY TCATGICCAT CATGGAGCAG CCGCTCGACC TGACCCACAT TGTGGSCACA GCAGAAGGCT TTAACCAAGC AGATAAGGCA ATAAACACCG CTGTAAAGTC GGGCAGGTGG GTGATGCTGA AGAATGTGCA TCTGGCCCCA GGGTGGCTGA TGCAGCTGGA GAAGAAGTTG CATTCCCTGC AGCCGCATGC CTGCTTCCGA CTCTTCCTCA CCATGGAGAT CAACCCCAAG GTGCCTGTGA ATCTGCTCCG TGCGGCCGC ATCTTTGTGT TCGAGCCACC GCCAGGGKTG GOGOTOGICA GOCTGAGCTG COTTCCCCC TITAAGGACT TGATTGCAAA GGTTCAGGCA GACGAGCAAT TIGGCATCTG CCTGGACAGC ACCTCCCCGG AGCAGACTGT GCCCTACCTC CAGGCTTTCC GGCCGATCG CCTGTTGGCC ATGCCCACA TGTTTGTTTC AACAAACCTT GAGGTGAAGC CCAACACTCC TGTCTTAATG TGCTCTGTGC CTGGTTATGA TGCCAGTGGA CATGITCGAGG ACCTTGCAGC CGAGCAGAAC ACGCAGATCA CTTCAATTGC AATCGCCTCT AAGGCCAACA TGCTGAGGAC GTTCAGCAGC ATTCCCGTCT CACGGATATG CAAGTCTCCC AACGAGCGTG CCCGCTTGTA CTTCCTGCTG GCCTGGTTTC ATGCGATCAT CCAAGAACGC TINCATACS CACCACTOGG GTGGTCAAAG AAGTATGAAT TTGGAGAGTC TGACCTGCGG

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50 45 4 \mathfrak{Z} 30 25 20 7 5 55 S GACCCTGAGC CACCTCAAGC GCACCGTGGA GAATATCAAG GATCCTTTGT TCAGGTTCTT CACACAGGGT ATTICAMATICO CAGATIGOCATI GCAGGIGGAGA GGAGTITIGTIG CAGTIGGGIGG AGTITICTICO GOCCOCOTOG ACAACGAGTT TGACCAGCGT CTGCTCAACA CCTTCCTGGA GCGCCTGTTC TCACCOGATA AGATICCOGTG GTCTGCACTA AAGACCTTAA TGGCCCAGTC CATTTATGGC TCANYTTOCG ATACGOTGGA CACGTGGCTG GATGACACGG CCAAGGGCAG GCAGAACATC 8 CAGCTICTAC GAGCGGGGTG TCGCAGTCTT GTGCACAGAG TAAACTTTTC TAGCTGCCCC CAAGCAGACA AACACCGAGA AGAAGGCCAG TGTGGTAACC TTACCTGTCT ACCTGAACTT CTOGTECETO GAGGAGETET GECTGGAAGT CAACGTEACC ACCTEACAGG GEGECACCET ACTOGOCAGOT GCATICTOGTG GOGOCIAAGGA GCTAAAGAAC ATCCACGTGT GCCTGGGTGG GACCGICATC CAGTGGGIST CCGACTICAG CGAGAGGATC AAACAGCTGC AGAACATCTC CCAGCTAGTG AAAGGGATCT TGCCTCGGAG CTGGTCCCAC TACACGGTGC CTGCCGGCAT TOTOGTOCAG GTOTOCOAAG GAAAGAAGAA GCAGACCAAC TACTTOCOCA COCTGATCAA TCAGAGAGAA OTGAAGATOG GCGCAAAGCT OCTTCAGGAC OTTCGCCAGG ACCTTGCAGA CCCTGCCTGG ATGCGGACAC TGCACACCAC CGCGTCCAAC TGGCTGCACC TCATCCCCCA CGACCTGGCC TACGCAGAGA CTGAGAAGAA GACGAGGACA GACTCCACGT CCGACGGGG CGACACCCAG ACGCCCTCCT GGCTGGGCCT GCCCAACAAC GCCGAGAGAG TCCTCCTTAC ACAACCAGGA GTTTCGACAG TGAGTTTAAG CTGGCATGCA AGGTCGACGG ACATAAAGAC TTTCTGTAAT AGTGAAAGTT GGTATTTAAC ATTTATTCAT TTTTAAAATA TTTGGAAGGT CACCOGIGCA GACCICATOT TCACOGIGGA CITOGAAAIT GCTACAAAGG AGGAICOTOG CAAGCIGICA CIGICCAAIG CCAICICAAC CGCCCTICCC CIGACGCAGC IGCGCIGGGI OCTOTICOTO COTGAGGOOT ACATCACIGO CACCAGGOAG TATOTOGOCO AGGOCAACAG GTGGACATGA TCAGTAAAAT GCTGAAGATG CAGATGTTGG AGGATGAGGA GGANATANAA CACTAAGCAT GANANAAAAA AANAAACTTA CAANCCNCAA GOTGGANATT GGANGGATAC CAGGAGGTAT TTGGGAAGGC CAATGGCGTG GAAAAGAAAG TOGTTOGTCT ACCITICUCAG TCACGGGITT GAAACTICAA GGGGCCACGT GCAACAACAA GAGGITOGAG GAAGCIGAAT GGAATCIGAC 2580 2520 2460 2400 2040 3360 3300 3240 3060 3000 2820 2700 2640 2280 2220

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INFORMATION FOR SEQ ID NO: 64:

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TEATEGCACE AAGGAGGTGC CCATGAACCC AGTGAAGATA TATCAAGTGT GTGACATCCC

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8 3 6 ઝ ಶ 23 5 5 S 8 GACATTCCTA TCACATATCC TACTACTGCC CCAGAAATTG CAGTTCCTGA GCTGGATGGA COTGAGITICT GOOTGCAGCG ACTGAAGGAG GAATATCAGT CCCTTATCCG GTATGTGGAG CTAGTTGTTA CTAAGTANTG CAGTAGCATT NTGGGGAAGA ACA ATACTAMITT CCIGIGCATC ACACTIAACT CATCTAACTG TICCCCGGAC ANCCICCACT TOGGGCCAGG AATGTGCCCA AATTTGGACT AGCTCATCTC ATGGCTCTGG GGCTGGGTCC AAGACAGCAA AGATOTACAG GOGTOGCAAA ATATGCCTGA COGATCATTT CAAACCTTTG TOGTTTOGAA AATGCTOGTA TATCCATGAC CTCCTGAAAT ATGAGTTTGA CATCGAGTTT AACAACAAGA ATOCTGACAA CGATTOOTTC CGACTGGAGT CCAACAAGGA AGGAACTCGG AGCTACAGGC AACACCACTT CCGCGTTTCT CTTGCGCCCT GGTCCAAGAT GGCGGATGAA GOTOTTACAG CAACCCAACG TOTCATOTTO CCATAGTAAA GATGACGGCG COTTGAGGTA CTCGATGCAC TCACAAGCGG GTAACTAGGT GACAAGAAAA CAAAGATCTT ATTCAAAAGA AGGIGATITT AATGATAGGT GICATATATA GGACGGATAA TCTGTTTACA TICTGTTCTT GATTICCCTOG AAACATOCCA CCCGGCATAG CCCTCAACAA GAAGAGGAAA ATACCATTIT 3 ATGGSTOOCA GTGGAAATCC CTGATCTGAT TCAGAAGGGC GTCATCCAAC ACAAAGAGAA OCCACOCOAC GIGITIGIGIC TOAGATOCOG GIOCIDAAGA CIAACOCOG ACCOCOAGAT TARATGGAAG GCCCAGCTGC GCTGTGCTCT CARTAAGAGC AGAGAATTCA ACCTGATGTA TAAGGCCTGG GCTGTAGAGA CAGGGAAGTA CCAGGAAGGG GTGGATGACC CTGACCCAGG GGCACGAGGT GGCCTCTACC CTGGGCTCAT CTGGCTACAC AGGGACTCTA AACGCTTCCA INFORMATION FOR SEQ ID NO: 65: E (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 65: SEQUENCE CHARACTERISTICS: TGAAGAATCA AGCCACTGAG GCAGGGCAGA GGGACCTTTG ATAGGCTACG (C) STRANDEDNESS: do: (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 1541 base pairs STRANDEDNESS: double 660 600 540 240 840 780 480 420 360 300 180 120 240 180 720 120 60 60

Ξ SEQUENCE CHARACTERISTICS:

(A) LENGTH: 883 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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SEQUENCE DESCRIPTION: SEQ ID NO: 64:

(D) TOPOLOGY: linear

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	TCAGCCCCAG GGCTCGATCA TTAACCCAGG ATCCACAGGG TCTGCTCCCT GGGATGAGAA	360	CCCTTCTIGTG TCTGCGTGAC CTCACCCAGC CTAGGAGGGA GGTGCATTCA GGGTAGATTT
~	GGATAATGAT GTGGATGAAG AAGATGAGGA AGATGAGCTG GATCAGTCGC AGCACCATGT	420	
-	TCCCATCCAG GACACCTTCC CCTTCCTGAA CATCAATGGT TCTCCCATGG CGCCAGCCAG	480	
	TOTOGOCAAT TOCAOTOTOG GCAACTOCAG CCCGGAGOCA OTOTOGCCCA AAACTGAACC	540	AZZOTOWIZZO OTAZZERZAM INCERNIZARA MININGOCIAL BAPTARABANING ANOVOTOM
01	CCTGGAGATG GAAGTACCC AGGCACTAT ACAGCCCTTC TATAGCTCTC CAGAACTGTG	600 10	
	GATCAGETET CTCCCAATGA CTGACCTGGA CATCAAGTTT CAGTACCOTG GGAAGGAGTA	999	GANCIAGAIS CLAGARAGEC 163CTIGIC CCCATGIGGS AGCCTGICC TCAGCCCTCT
2	COGOCAGACE ATCACCOTCA GCACCCTCA GOCCTGCCGA CTCTTCTATG GCGACCTGGG	720	
<u>C</u>	TCCCATGCCT GACCAGGAGG AGCTCTTTGG TCCCGTCAGA CTGGAGCAGG TCAAATTCCC	084	AININIUGAS AGAGITUGAS AGININIUGA GAMINITITI GGAAAGAGI TUGTUTIGG AAUGGUSAGU IQOLAAGAGU IQOLAAGAGU AAAGAGU AAAGAGU AAAAAAAAAAAAAAAA
	AGGTECTICAG CATATTACCA ATGACAAGCA CAAGCTOTTE ACTAGCAAGC TOCTOGAGGT	840	ANTOLOGIS INCOMESTI PARTOLISTA INCOMESTI INCOMESTI
20	CATOGACAGA GGACTGATCC TGGAGGTCAG GGGTCATGCC ATTINTGCCA TCAGGCTGTG	900 20	AAAGGICIAC
	CCAGTGCAAG GTGTACTGGT CTGGGCCATG TGCCCCATCA CTTGTTGCTC CCAACCTGAT	096	to concensor
ç	TGAGAGACAA AAGAAGGTCA AGCTATTTTG TCTGGAAACA TTCCTTAGGG ATCTCATTGC	1020	
3	CCACCAGAAA GGACAGATAG AGAAGCAGCC ACCOTTTGAG ATCTACTTAT GCTTTTGGGA	1080	(2) INFORMATION FOR SEQ ID NO: 67:
	AGAATGGCCA GATGGGAAAC CATTGGAAAG GAAACTCATC TTGGTTCAGG TCATTCCAGT	1140	(i) SEQUENCE CHARACTERISTICS:
30	AGIGGCTCGG ATGATCTACG AGATGTTTTC TGGTGATTTC ACACGATCCT TTGATAGTGG	1200 30	(A) LEGALIST OS DAME PAIRS (B) TYPES INCIDED CALL (C) CTERNAME PRINCE CO. ALIA D.
	CAGTOTOCOC CTOCAGATOT CAACOCCAGA CATCAAGGAF AACATOOTTO CTCAGCTGAA	1260	(c) SirAwbaracass: compact(D) TOPOLOGY: Linear
35	GCAGCTIGTAC CGCATCCTTC AAACCCAGGA GAGCTGGCAG CCCATGCAGC CCACCCCCAG	1320	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 67:
4	CATGCAACTG CCCCCTGCCC TGCCTCCCCA GTAATTGTGA ATGCCATCTT CTTCCTTCTC	1380	TTAAGGAATT CGGCKGATC CCGGCAAGTA ACATGACTAA AAAGAAGCGG GAGAATCTGG
	TITITIADA TATIGIACAT ARGGATTITT ITATICITIA CATITAACCA CCITITAAAL	1440	OCCITCATOR AGAGATICAN GOCTAGAGO AGAACTICIC CCAGITOTCOG AGAGACCTGG
40	CICTOTITIC TOTCACAGIG TIMGAGGITT GIGATICICC AMAINIOCCT AGAITTAAAG	1500 40	наяссетам стеснанете сисмассвая местамаесе манавсемая мастесетва
	CTGATTTAAT TTATGGAAAA AAAAAAAAA AAAAAAAAA	1541	AGNAGGAGNA ANACHGCTN ATGNACANAG CCTCCNACTA CGNGNAGGNA CTGNAGTTTC
45		45	TICOSCIAGA GAACCIGIAG AACATIGCTICC TCTCTIGTIGGC CATCTITTATIC CTCCTGACGC
	(2) INFORMATION FOR SEQ ID NO: 66:		TOSTETATISC CTACTIGAACC ATGTGAACCT GGCACTTCCC CACAACCAAC ACAGGCTTCC
Š	(i) SEQ		ACTITIONICCE TITOSTICAGOA TCAAGCHOOC ACTITICAAGCC TCAATAGGGAC CAAGGTTGCTG
ಗ	(B)		SESTING CICCARCCI ASIGNICANO CANSCINCI NECESCULA GEORIGA
	(C) STRANDEDMESS: double (D) TOPOLGGY: linear		CCTIGACTIG CTGGGGGTT CCGGGTCTCC AGAGGACAT GGGCTGGTC CCTCCCTTAG
55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:	55	CCCANGGGAG AGGCAATAAN GAACACAAAG CTGAAAAAAA AAAAAAAAA AACTCGTAGG
	AGANANTGAN TOTTHGANGG TECCTECCEA GOCGGGACAG AGTOTTTGCT COCCCTGGAG	09	GOGGOCCOST ACCCANTOGC CCTINICOTO
9	AAGGCTCTGC TCAGCCCTTGA GAGTCCCTTC CTGCCCCACC GATACTGGCA CTTTAAAAAG	120 60	
;			

AGCOGTIGATIC ATROGGGGCCC CGGGCTCGGG CAAGGGGCACC GTGTCGTCGC GCATCACTAC CACAGIGATT AACCIGAATG TOCCCITTGA GGICATIAAA CAACGCCITA CIGCIGGCIG CTOCTAGECG GCCGGCGCAG GCTGCCGAGC GGGTGAGCGC GCAGGCCAGG CCAAAGCCCT GTTCAGTTAA TAAGTGGTTG ATAAAGTTTC CATATTTTTC TGGAAAAGTT AAAAAAAGTT ACCTITAAAA CATCIGITAG AGCAAAATTA AAAGAGCATT TOGTAGTAAT CTAACTITTT GCTGCTTTTC CTAAGACTTC TAGTATGTAT GAATTCTTTG AMAGTATAT TACTTTTATT TOCTITICCTA CARACTARAG TICCACARAG ARGCCAGRAR GCTICAGTTA CICCATGROG AAAAGGGGTG CTGGAAACAT TCTCCGGAAC AGAAACCAAC AAGATTTGGC CCTATGTATA TATCAAQAGA CTAAAQGCTT ATGAAGACCA AACAAAGCCA GTCCTGGAAT ATTACCAGAA CATTGATGAC CTGACTGGGG AGCCTCTCAT TCAGCGTGAG GATGATAAAC CAGAGACGGT GATTCATCCC GCCAGTOGCC GAGTCTATAA CATTGAATTC AACCCTCCCA AAACTGTOGG ACATGROATT TOGAGAAAAT ACGTAATCAG AAATTTTGTGC ATAGATTGAT GCCAAAAAAG COTTIGAAAT CATCTAGIGT GTIGTATGCA GTTATCCTCA AAAACATCAG CGATGTCTGA ACACTTOGAG CTGAAGCACC TCTCCAGCGG GGACCTGCTC CGGGACAACA TGCTGCGGGG TATGTTAACC ATATATGCTG TATTTATTTT GTCGTTAAGC ATACTTTCAG TTTACTCAGA ACATTICCAG CATIGIGGAA CAIGGIGAGA CACIATATAA AATICCAGAA AGAAAGCAAC TOGATTTACA GATTTATTOT GAGACACAAA TTCACTOCTG CCTTTACACT AAGAAATOTA ATTTTCAATT TOCTATAAAG ATGTATCAAT TAGCATATAG AAAAATATTA CTTTAAGATG TTATTTTGGA TACTAAGGAT GTGCCAAATG ATTCGGATAC TAAGATGCAT GTAACTATTA ATAGTAAGAT GGGCAAACCT CCTAGTCCTT GCATTTAGAA CCAAGGACAC TICCACAGGC AGAAGCCCTA GATAGAGCTI ATCAGATCGA COGCIOCCC TICATGAGCT GAAAAATCIC ACCCAGIATA GCIGGCIGIT GOCGTOTTAG CCAAGGCTTT CATTGACCAA GGGAAACTCA TCCCAGATGA COGTOCOGOC CTCACTICTGC GOCCATOGGG GCGTCCGCGC GGCTGCTGCG SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 1751 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 291 PCT/US98/04493 1080 1020 480 420 360 8 180 120 720 660 600 900 780 23 20 330 15 5 \$ 6 Š 50 WO 98/39448 TRAGTOCTTT TACHAGCACC ARAGTTCART GRATTTTCAR CRARATGTRA TTARAGTCTA TITIGATICCA AATGIGIGAT CIGCCCIGAT AAATAACAAG TIATNOTACC AICTCCCCCG TOTTTTCAGT TATGACTCAG GITTAAGAAAT GIGITTITAGG ATCTACTIGC CAGCTAAAAA TIGAAAACAA AGATCITGGAC AACAAAACAG CCAAAGGTIGG GGGTCAAGAA GCCACGAGAT TATGTATTAA AATGTTTTTG AATTGTGAAA TATTAGAATA TIGTTACTAT CCANTAAAA AAAAAAAAA AAAAAAAAAC TCGAGGGGG GCCCGGTACC CAATTCTCCG CCTACTCAGC TGGACTTAAA AATAAAAA ATCOTOCADA GCATTADAGA DATCTTOTTA CTGCTADAGTO TTGCTGACCC AGGAACAACT CCAGGLAAAA CTGCAGAGAG CCCCAGTCTT CACCTCTGGT TGACCATGAG CTCTGTGTAA OCTOTOLOGI: GTACCTAGOT GTAGAATOCT ATGCACACGT GCCAGGTGTA GTGTGCATAT ATAMICAGAT GCAGTATCAC AGCTGTGTCA GACTCTAGTA CCAGTTGGGC AATCAAGGCA TIGACCCAAC ICAAAAICIC CAIGGGAAAA TACCIGICGA TACCCACAGI ATIGIIGAAA 5 NAATAGGNAG T 3 GCAGGAAGTG GATTOCAGAT TIGCTIGIGI CCICAGGIGA IGANGAGGG IGITITICCCC IGITIGICCIT TACATAGAGC AAAGAGAAAT TITCCAGAATT TCTARAATTC TGGAAAGAGA ATTITCCTGA TECTEACACT CATECTTECT CIECTAGAGT GICTEGTING CATGATCATE TECTACCIAG INFORMATION FOR SEQ ID NO: 69: INFORMATION FOR SEQ ID NO: 70: Ξ Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70: SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: ARGECTARGE CAGATTTRAG CTCTGRARAGE ATTCCRCARC ATRCRCACA SEQUENCE DESCRIPTION: SEQ ID NO: 69: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 508 base pairs (B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear (A) LENGTH: 245 base pairs STRANDEDNESS: double 292

> 240 180 120

480 420 360 300

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GATGGTTTT

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ACTIGITICC TITGAAAATA CCIGIGIACI GAGGGITAIG ATTIGIGICA AAAATIGACA

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AGAAATGTGT

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SEQUENCE CHARACTERISTICS:

IGGITITICI

1620 1560

1680

1751 1740 PCT/US98/04493

INFORMATION FOR SEQ ID NO: 68:

5		PCT/tJS98/04493	. ≯	WO 98/39448 PC	PCT/US98/04493	
•	293			294		
	GCATTICTIT CACTGATACA AGGAAAACTG CAGGGTTAAA AAAAAAAAA AAAAAAAAA	240		GICTITICCT GITAGAITT TAATAGCAGA ACTGIATGAC AAGTTIAGGT GATCCTAGCA	540	
	NCNCO	245		TATOTTANAT TCANATTANT GTANAACAGA TTANCANCAA CANAGANACT GTCTATTTGA	009	
5			5	GIGANGTCAT GCTTTCTATT ATAATAACTT GGCTTCGGTT ATCCATCAAA TGCACACTTA	099	
	: : : : : : : : : : : : : : : : : : : :			TACTOTTATC TGATTOTTTA TAATAAAGAA TACTOTACTT ATAAAAAAA AAA	713	•
	(2) INFORMATION FOR SEQ ID NO: /I:		9			
2	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 361 base pairs		2			
				(2) INFORMATION FOR SEQ ID NO: 73:		
:	9 9		ž	(1) SEQUENCE CHARACTERISTICS:		
2	(x1) SEQUENCE DESCRIPTION: SEQ 1D NO: 71:		2			
	ATGITCCTCA TGAGGATGCA CITGTGCTTC TGCAAGTAIT GCTGCAGCTT CATAGTGACT	09		(D) TOPOLOGY: Linear		
20	CCCACCAGCA CCAGCAATAC AGCTAGCTAC CTGTGGCCTT GANTOTCAGC CAGCATGGCT	120	70	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:		
	GOSAGAGGGA GCAGCTGGGC ATGTACCCTA AATGCTGTTA CCAGGGAAGG ACTCCCAGAG	180		GAAAGTCAGA GCTGTCCAAT CCCTCAGCAC CTTTTAGATT TGCTCCAAAT TAGAAACGTG	09	
;	TGAAGACAAG TAGGGACTTC CTGCAGAGGT GGTACATGTG CTCTCTGTAT CCATACTTTT	240	ý	GOGACTATOT GITCTGGGCA ATCACAGGTC TGGAAAATGG CTCTGCAGGC TCTTGATAGT	120	
3	TITITITITI TITIDAGATA GAGTITICACC CITOTIGOCC TOGCIGGAGT GCAATGGTGC	300	3	GAGACAGTGG TCATCTTACC AGACATGCAT CTGATTTTAA GCCTCAGGCT AATCCACAAT	180	
	GATCTCAGCT CACTGCAACC TCTCTGCCTC CCGGGTTCAA GTGATTCTCC TGCCTCAGCC	360		GCTCGGCCAT GCCTATGAIT AACAAACAAA AGCAAAATCT GCTTTTATAG TTTAGGAAAC	240	
3	· ·	361	30	CISGARAGAA CAGIATITITI CAGCATICITI GGARAAAGCA GITCIGCAITI TITAAATIGG	300	
				gactgcagaa otgactotet atagttotga aatacaaaaa atgotatott tgatcagaaa	360	
3			35	AGGAAGCCCG TGCCTGGCAC TTGGAAAGAT ACTGAGCATC ATAACCCTAA TGAGAAAATG	420	
દ	(2) INFORMATION FOR SEQ ID NO: 72:		3	THEGOTICIET GARIETTARC TACARATICAG GITHEGRANG CATATIGACAC CCTITIGICAA	_480	
	(1) SEQUENCE CHARACTERISTICS: (A) LENOTH: 713 base pairs			ACTIVACCITIC ACTIVICAGAGGA CCTIGTOCTICA TAGAAGAATA TOCTITTAAAA GTATICAATITT		
9			40	TCCACAOTCG ATGATGGAGA AAAGTTCATT TGCACCAGAA TGCTGATAGT CACAATACAC	900_	
	(D) TOPOLOGY: linear			AGCETGACAT ATATAACAAT ACAGTTTTCT GTAAACAGAA GTTCTTCCTC TTCCAATTCA	. 099	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:		3	CCACTCAGTC ACACCATAAA TATTGCATGT TTCACTTTAG AAACTGATTC ATTTTAGAAA	720	
45	AGGATCACAC AATAGAGAAC ACTGTAGTAA CATTTCGGTC TOCTCACAAG ACCCAGAACA	09	.	GCAGATCTGG ATTAITTTGC AGGTAGAAA TGAAGGCTAT TTCTGGCATT CTTGCTCAAA	780	
	ITIGAICAGIT ITIGITIGITIG GITTAITAIT ITICIGITAA AAAITIGIGA AAAGITIGIT	120		ANGICAATAT ATGTACATTA AGTATAAAAA AGGGTCTCTT TCACCTCTTT TGTTTTGTAG	840	
20	THACTAGAT GATATITHAA TAGCIGOGAG IGCTITIGGAA CHAIAAAGAT GICACHACTT	180	20	CATTGGCTAC ATAACTCGTG CC	 862	
3	AACACACATA CCTTATGTTT TGTTTTACAC TCAGTATAAA TCAGGAGAAG	240				
	TTAGCCAACC ATCTAGCATT TAGAATCCTC TTTTTTATTG TCTTCTAAGG ATATGGATGT	300				
55	TCCCATAACA GCAACAAAC AGCAACAAAA ACATTTCATA AATATCACTT GATAGACTOT	360	55	(2) INFORMATION FOR SEQ ID NO: 74:	-	
	AAGCACCIGC TIAACITIIGT GTCCCAAAIA TITAGIGIGT AIAIATATA ATATATATAC	420		(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 4602 base pairs		
9	ACACACACA ACATATÁTAT TCAACAAATA AAGCAAAATA TAACATGCAT TTCACATTT	480	9			

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296

30 23 8 15 8 35 5 3 55 50 8 AAAATCTAAA GCATTTAAAA TCTAGTGAAA TAACTGAAGG GCCTGCTCTT TCCATTGTGG GAAGACTTOG GGCATGGGAC AGCTCAGACT TTGTATTTAA AAGTTAAAAA GGACAAAAAA TETTTOTOGO AAGCGGCAGC AGTGGAGGCA CTGAAGGGCT GGTTATGAAC TCAGATATAC TOGGCAGATO COCCTITICCA ACCIGIAACT CIGATGIGCI CIGGATCAGC TITTAACTIT GCTCTTAAAG TGAGGGTTAT TCTCATACTC GGTTCCAGCC ATCAGCAGAC TTCCTGCTCA TOCCCTTATA AAATTGATGT TGTCTTTACC AGAAAGGTAG ACAAAAAAAGA AGCAGCAGCA ATCACAGCAC ACACATACAT ACACCCICCA CCTCCCCATC CCCTGTTCTC CCTCTGTTGC TOGOTOCTAC CACAGAGGTT CTGATTGAAG ATTCAGACTC TOCCOGACCT TAGTGGACAG GITCAGAATC TAAATTACAG ATAGATGATT GITTCITGIG AATTIGITIC TITTCCITTT AUGCTOTTIG AGAITTITIAC AGIGTIGAAA CITAAGAAIT TIGAGAGGGI GAGGAGGGIT TAGCCTAGIG CTITTTIGGA AGCCTITTIA GGGAAGAATG TTAGGTICAT GGTAACTAGT CATACCTCTC CACTOCCTTT TAAAGAAAGC TOGTCCTCAG CACTAACAAA ATCACTACAA OCCIOCIACO COCIOCOGCA CACITATOTI CAAATACCAT AGAATICTAA ICTOTGAAAT TAATCATATA TTACTOTCTT CTAAATCCCT TCTCCTCCTC TACTGCTGCC CTATGGTTCT OTCAGAAATG CATGATGGCT CTTGGAAAGA ATGACGTTTT GCTGGAAAAA AAAAAAARAA TAATAATTOC TTTGGCTTTC ACCTAAAATT CTGGGCATCA CAATTTCCTT GGGATAGAG CCCCTTTAGT AACACTICTG AAGAGGAAAA ACTICAATAG CCAAAGITAA TAATCCIATA TAGTGAAGGA ACAAAGTCTA TGAGTCCTAA AATTTTAAGT CAAAGAAAAC TGCTCTGTTT TOTTICTIOC TITICTITAT CAGTICATIC CAGCICCCIG TIAGIGAAGG ACACIGCIGT TITTIONCCC TACCATINCC TRACATINCC CTIGGGGCCC ANCICIOGCI CCTIGCITT AGAGGAGTAA CAAAATGTCA TITCTGAAAG AGGCTTACTT TATACCAACT AGTGTCAGCA CMSTTTGTGT TICACAAACA TOGCTTATCA ATTTTTTCAA AGAATTCTTT TITCCCAAAA ATTECTATES TOTOTOCCTS AGAAGSTAGA ATSGAAGGST TASSSTATTS ATTGTTATT ACAAGGTCAT ACCOCCAGAA GCCCCAAATC CTATTTTOGC TCATCTTCAG GTAAAGAGTA TIGROTIGGG GAARAGATIG CTIATIOCIG TICACIOGAG AGAAAAGGIA GIGTTITIOT ANTARAGATA GARGGGATGT AGRATIATGCT TICCTCCCAA CATGGITTGG AGTCGACTIT TITGGGATGC CAGGGAACAG AGAGTGAGAC ACCTACAATC ACCAGTCTCA AATGCGCTAT CICCICICAG GICCCITITA CACTITITGA CIAACIAGCA ICIATATICC ACACTIAGCI GGTATATTGA CTAGATTTGA AAATACAAGA TIGATTAGAT GAATCTACAA AAAAGTTGTC TGITTCTTTT CAGAGIGTIG CAGAITTGCC ATTICIOCAT AATAIGGGGA TAGAAAATGG 2160 2040 1980 1920 1860 1800 1740 2880 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2220 3060 3240 3180 3120 3000 2940 3480 3420 3360 3300

20 5 5 35 30 23 45 6 S 8 55 CCCAGGGGGG GKGGGGAGCA GCGCCGARGC CGCCCCCCCC GCCTCCGCCG CCTAGGACTA GATTETECCC TECCTEAGGA AGITTICEATT GETGCATETA GACCTAGECG GGGCTGGCGT CTATGCATAG AAAAAGTAAC AACTGATAAA GATCCCAAGG AAGAAAAAGA GGAAGAAGAG GCCCCCTGAG GCGGTCGCGG CGGCGCCTGC AGGGACCACT AGCAGCCGCG TGCTGAGGGG GCAGCAGGAC CAGGAAGGGG GAGAGGCGGC CAAGGCGGCT CCGGAGGACC CGCAACAACG вороствова алекаленна ссессантнее воскальнее алланистал влессества GATOTTOGAG AAGAGCATCA GTCTCCAGGT GGCATTAGTA GTGAAGAGGA AGAGGAGGAG AAGACCOGTT CATTGCAGCT CATTTGCAAG TCAGAACCAA ATACAGACCA ACTTGATTAT AGTAGTAGGA CATCTGTTTC TCGCCATCGT GATACAGAGA ACACCCGAAG CTCTCGGTCC GEAGEAGECE AGGECGEGAA GIECCEGIET CEAGITEAGG GEAAGAAGAG ICCGEGAETE AGGCCGAGTA TCCCCGGCGG CGAGGAGCAG CCCCAGGGGC AGGCCTCCCG ACGTCCCGG даглеваале саловесава есоствестне соссветами сластвтоге освесавлал TACCAGCATT TOCTGAAGAA GAAATATOTA TOTOCCCATC OCTOCTOTOG ACGACTOTTO GAGAATGAAA TITAGAGAGGA TGAGGAACCT CCAAGGAAGA GAGGAAGAAG ACGAAAAGAT ACCTACAAAC CCCACTTAGA AAGGGAAAACC CCAAAGCCAC GGAGAAAATC AGGGAAGGTA GAAGAAGAGA TOTTAATCAG TGAAGAGGAG ATACCATTCA AAGATGATCC AAGAGATGAG CTCTTHATTG GCACATGAAG AAACATGATG CAGACTCCTT CTACCAGTTT TCTTGCAATA CACACTOSCG AGAAGCATTA CAATGTGAGA TCTGTGGATT TACTTGTCGA CAAAAGGCAT TOTGAATATT GTGCTCGGGC CTTCAAGAGT TCCCACAATC TGGCAGTGCA CCGGATGATT aaagaagaga aogagaagaa ggaaattaaa otogaaotag aostogaost gaaagaagag AGGETTEAGA AGEAAETTET GEGACATGEE AAACATEATA CAGATEAAAG GGATTATATE ATATETTOGG CACTAACCCA GAGTCCCTGA CGCAGCCTTC AGATGGTCAG GGTCTTCCTC CTOHOGTOCT GATTOCAGAA GCTCTGGCTG CCAATGCAGG CGCCCTCATC ACCAGCACAG TOTOTOGOCAA AAAATTTOAG AAGAAGGACA GOGTAGTGGC ACACAAGGCA AAAAGCCACO GGATGICAAA GICATACIGC AGIGGGACGG AACGGGIGAG CCIGAIGGCI GAIGGGAAG TICTICCIGA GCCCITGGGA AACICAACCI CIGGAGAGIG CCIACIGITA GAAGCIGAAC GATGTGGAAC TGTCCTTGCC CATCCTCGCT ATTTGCAGCA CCACATTAAA CACCITIACC CAAAAGGAGA AAAAAGCCIC CAAICCAGIA IGICCGIIGI 1320 1260 1200 1140 1080 1020 1500 1380 840 780 720 660 600 540 180 120 1680 1620 1560 1440 960 900 480 420

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(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

š	WO 98/39448 P.	PCT/US98/04493	ž	WO 98/39448
	297	-		298
	TITITIGICAC ACTIVITECTI IGIETECGIA AATITEATIT GEAGIGGITA GICATEAGAI	3540		GCCAAAAGAT ATTTGACGGT TTCCAAAATT CAGATTCTGC CTCTGCGGAT AAATATTTGC
	ATTITIAGCCA CCTACACAAA AGCAAACTGC ATTITIAAAA AICTITICTGA GATGGGAGAA	3600		CACAMITANG TANGFOCKET CACCACTOTS ANGESTORICA CACAMICATTY TOACACTORS
2	AATGTATTCT CCTTTCCTAT ACGCCTCTCC CAACAAAAA ACAACTAGTT AGTTCTACTA	3660	5	TPAGEACTICA ACTIVITICATE GANCTAGGAT GAICTICATIVC COCINCIGGAT GAACATOCIC
	ATTAGABACT TECTETACIT TITETITICT TITAGGGGTC ANGANCECTE TITATAGGTA	3720		TIGATIGATICA GESTICCAGE ARESTACTIT GAAGGGAACA ATCAGATICA AAAGCTICTIG
9	CCAITIGCCT ACAADAAATT ATTGCAGCAG TTTGCAATAC TAAAATATTT TTTAIAGACT	3780	2	GOTTHTIATT TARABITATE CHESTOACHT CYGAGTRACCC GCCGCTUCAC ACCCTUAGET
2	THARAITITT CCTTTTGATA AAGGGATGCT GCATAGTAGA GTTGGTGTAA TTAAACTATC	3840	2	CARGOCITATA DECENSOR DE CONCATATIVO ATTITICATOR
	TCAGCOGITT CCCTGCTITC CCTTCTGCTC CATAGGCTC ATTGTCCTTC CAGGGAGCTC	3900		THE TRANSCOTT CACCIOCARA GICACICITY TRANSCORGE GAACCAAAT GETGANGATA
15	TITTARICIT ARAGITCIAC ATTICATECT CITAGICAA ITCIGITACC TITTIARINA	3960	15	TAINGAKTITT AUGUNTAGC ACAGITCANC CCCAACCCTA GYCTYCGAAA IGTIAARATT
	CICITCCCAC TOCATATTC CATCITGAAT TOCTGGTTCT AMATTCTGAA ACTGTAGTTG	4020		TGAGAAARTCE AGAAAARGCA TECARAART TACAGAARTIC AAARATIGGA AAAGGARGGIG
6	AGATACAGCT ATTTAATATT TCTGGGAGAT GTGCATCCCT CTTCTTTGTG GTTGCCCAAG	4080	70	TOTALTICIA COCCARGOTOC COTOTICOS ACCACCACOS TACCATATA
97	STIGHTHISC STANCHSAGA CTCCTTGATA TGCTTCAGAG AATTTAGGCA AACACTGGCC	4140	1	HENETRANCIA GESCHAMENA CONCRAGECA CONCINERANG CACHOOOCE THORIGITALS
	ATGCCCTIGG GAGTACTIGGG AUTABANTAA AAMTATCGAG GTATAGACTA GCATCCACAT	4200		ACABACHTER COCHEMBAN CHARACHTCH WINTHWATTH BARARANG COCHEMBANA CHARACHTER AND AND AND AND AND AND AND AND AND AND
25	AGAGCACTIG AACCICCITI GIACCIGITI GOGGAAAAAG TAIAAIGAGI GIACTACCAA	4260	25	PROTECTION OF THE PROPERTY OF
	TCHALCHAG ATTATTATAG TCTGGTTGTT TGAAATACCA TTTTTTTCTC CTTTTGTGTT	4320		AMARITANCA MACACAMBAR MINAMBARA MARITANCANA ARABITANCAN MARITAN MARITANCAN MA
	TITICCCACITI TCCANTGIAC TCAAGAAAT TGAACAAATG TAATGGATCA AITTAAAATA	4380	9	nivamino mmemmino persenta sessentano sessentano mandelessa
30	TITIATHET TANAGECTT TITIGECTOT TOTANISTIC AGREECTTE TECTITICATE	4440	3	TICCITITI TICITITI CETTO TO TO THE TOTAL CITICAGE
	ggagagacag stagttacct gaatataggt tgaaaagstt atstaaaag aaattataat	4500		MANAGARIA MANAGARIA MANAGARIA CECENORIA MENTE
35	AAAAGGATA CTTIGCTITT CAAATCTITG ITTICICITA TICIAGGIAA GGCATATIAA	4560	35	
	aaataaatat gtaaagaaga aaaataaaag ttgtcttcat go	4602		(2) INFORMATION FOR SEQ ID NO: 76:
9			40	
	(2) INFORMATION FOR SEQ ID NO: 75:			(B) TYPE: INUCIAIC ACLO (C) STRANDENESS: double (D) TOPOLOSY: linear
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1255 base pairs		45	
!	(B) TYPE: nucleic acid			GGCACGAGAG AAANGTITIGA TICTCTITICC TATTITAAGG GANCTICTICT CTIGTIGAIG
				THE STATE AND APPEARED AND ACTORISED OF ACTORISED THE STATE THE STATE OF THE STATE AND ACTORISES.
20	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 75:		20	CACAGCIATO CANCIATICA AAGTGATGAT CIOTIGGGATA GITTTAATGA GGTCACAAAC
	COCCCCCCG GCCGCGGGT TTCTCTAACA AATAAACAGA ACCCGCACTG CCCAAGGGAG	09		CANACACTAG ATGTAAAGAG AATGATGAAA ACCTGGACCC TGCAGAAAGG ATTTCCTTTA
¥	COTTOCCACT TTCAAAGTGG TCCCCTGGGG GAGCTCAGCC TCATCCTGAT GATGCTGCCA	120	55	GTGACTGITC AAAAGAAAGG AAAGGAACTT TITATACAAC AAGAGAGATT CITTITAAAT
ર	AGGCGCACIT TITATITITA ITITATITIT AITITITITI TAGCARCCIT TIGGGGCTIC	180	•	ATGARGCCTG ANATICAGCC TTCAGATACA AGGTACATGC CCTCTTTCTT TTCATGCCAT
	ACTICTICAGAS CCAGTITITIA AGGGACACCA GAGCCGCAGC CTGCTCTGAT TCTATGGCTT	240		CICTITUGCA CICTCAGGIG GAARIATITI GAAGIGITIT ATAATCATAA GITCITIGGA
9	GGITGTIACT ATAAGAGIAA TIGCCTAACT TGAITHTITCA TCTCTITIAAC CAAACTIGIG	300	9	

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299

AACCTAACAA GATTATCCCT TCCTAAGAAT ACTTAACCTT CCTACCAAAT TAAAA

475

2 INFORMATION FOR SEQ ID NO: 77:

Ξ SEQUENCE CHARACTERISTICS:

LENGTH: 465 base pairs

9 9 9 9 STRANDEDNESS: double TYPE: nucleic acid

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TOPOLOGY: linear

(X) SEQUENCE DESCRIPTION: SEQ ID NO: 77

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25 20 Trerecce GIGGGCGTCC CAGCOGTOCT GCCATOGCGT GGCGGCGCG CGAACCRGCG TCGGGGCTCG TICICICIOS TETICGACIO CACCOCACIO GOGOTIGACO CIGACIOCO CIAGICAGOI AGECCEGEGG CCGAGGGGCC GCGCCCTGGG TCGCCCTGGT GGTCGTGGGG COTOGOCOCC COGCOGAGAM CTCGARGOCT KCGCGCCCGA CACGCGCTTC AGAGTGOCCG CTTCGGCGAC AGCTCGCCCA AGGAGGGCGC GCATGGCCTG COGTIGITADA CATOGAGIAC GIGGACCCGC AGACCAACCI GACGGIGIGG TOCTOGECET GECCETOTICE GTGCCCGGGG CCCGGGGCCG GGCTCTCGAG CGCCGIGITIC 180 420 360 300 240 120

2 INFORMATION FOR SEQ ID NO: 78:

႘

30

GCTGCACCTT TCAAGGACAA AGTGCTGGTG GCGGCGCNGA ANGAA

465

Ξ SEQUENCE CHARACTERISTICS: LENGTH: 1907 base pairs

TOPOLOGY: linear

999 TYPE: nucleic acid STRANDEDNESS: double

6

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 78:

3 8 S 8 GGAAGAAATG AGTGGCTTAT ACTCTCCTGT CAGTGAGGAT AGCACAGTGC CTCAATTTGA ACATGCAGCC CAACTACAGA TICITATGGA ATTCCTCAAG GTTGCAAGAA GAAATAAGAG GTCTCGTATC TCAGATGACA GTCGAACTGC AAGCCAGTTG GATGAATTTC AGGAATGCTT ACGAMAACGA CTTACTGCTC ATTTTGAMGA CTTGGAGCAG TOTTACTTTT CTACAAGGAI TITICAGIGGC AGITICICAGA CAAAGAAACA GCCITGGIAT AAIAGCACGI TAGCATCAAG GICCAAGITT ACICGATATA ATTCAGTACG ACCITTAGCC ACATTGTCAT ATGCTAGTG AGCTCCTTCT CCATCACACA GTAGTATTAT TGATTCCACA GAATACAGCC AACCTCCAGG AGAGCAACTO GAACAGATCC AGAAGGAGCT AAGTSTTTTG GAAGAGGATA TTAAGAGAGT **4BO** 240 180 120 420 360 300 S

> 20 5 30 25 2 25 8 ઝ S TOCAGTOGAT ATTCATTACC CIGAGAATGA AATGACCTGC AATTCGAAAA TCAGCTGTAT TETETATAAT GOTTECAGTA TAGTETETAG TATTGAATTT GACCGGGATT GTGACTATTT TIGIGGAAGI GAAAATAACI CICICIACCI GIACIAIAAA GGACITICIA AGACITIGCI CAAGGSTCAT ATCAATGAAA AAAACTTTGT AGGCCTGGCT TCCAATGGAG ATTATATAGC CTCAACAGAC AGTCAGCTAA AACTGTGGAA TGTAGGGAAA CCATACTGCC TACGTTCCTT TAMIGIGIGE IGIGITAMAN TEMSECCETE TICCAGATAC CATTIOGETT TEGGETGIGE AAAAGTGAAG CIGIOGICIA CCAAICTAGA CAACICAGIG GCAAGCATIG AGGCAAAGGC TIGGAGIGIT GACITIAATI IGAIGGAICC TAAACICIIG GCIICAGGIT CIGAIGAIGC TITATOGGAT GGATTCACAG GACAGAGOTC AAAGOTCTAT CAGGAGCATG AGAAGAGOTG CAGTIGGAGI AGITACCATA AGAACCIGIT AGCIAGCAGI GATIAIGAAG GCACIGITAI TOCCATTOCT GGAGTTACAA AGAAGATTAA AGTCTATGAA TATGACACTG TCATCCAGGA CAACTCTCCA CCATCAATOT AACTCCATGG ACATTGCTGC TCTTGGTGGT GTTATCTAAT GATTOCTOCT AACAGICAGG GTACAATTAA GGIGCIAGAA TIGGIAIGAA AMATGMATTT GITMGTGCTG TGTGCTGGAG GGCACTACCA GATGGGGAGT AACTITIAAG TITGATACAG TCAAAAGTGT TCTCGACAAA GACCGAAAAG AAGATGATAC AGGACACCOT AAAGCAGICT CITATGCAAA GITIGTGAGT GGIGAGGAAA TIGTCICTGC AGATCACTOT GTCCACTACT ATGATCTTCG TAACACTAAA CAGCCAATCA TGGTATTCAA ANGANGTAGC ATATOTGAAC TATAATOTAA CAGTGAATAA TTTGTAAAGT TCGTATTTCC TITTOTGATA GOGAAACAAA TICTITIGAA TAAAAATAAA TAACAAAACA ATAAAAGITT ATTITITIET TITITICITY TECCICCITY ANGACCITYG GGACATIGGG AATACCCAGC AGAAAATGIC ATGIGATGIC TCTCCCCAAA GICATCATGG GITTITGGATT TGTTTTGAAT ANGICAAATT GIRCITGAIC CIGCIGAAAT ACAICIGCAG CIGACAAIGA GAGAAGAAAC ATTOAGCCAC AGITGAGCIT GGAAAGITITI TGTCAAATGC NGCAAGAGAI AACICTITITT CAACCTCTTT GGGAATTACA CATATCAATA TAAACAAAAT ATAAAGT GGGTTAACTC CCAATGTGCT 1500 1440 1380 1320 1200 1140 1080 1020 1560 1260 1907 1860 1800 1740 1680 1620 960 900 840 780 720 660 600 540

INFORMATION FOR SEQ ID NO: 79:

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Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: doub (A) LENGTH: 1168 base pairs STRANDEDNESS: double

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79: TOPOLOGY: linear

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TCAACTAAAG AAGATTTGCA TGCAGCGGTT CATTAAAAIC GATGGCAAGG ICCCAAACTGA	20		8
AMMOINTE CLUSTAINA IIIINOIMA ANAMANIA MANAMANA MANAMANA MANAMANA MANAMANA MANAMANA		GAGAAATGTT CAGGCTTGTA GGGATGGCAC ACTTATTAGT TCTGCCTGTC TGAAAGGTTC	
PORTERIOR CONTRAINED BUTTIVONED CARCACTOR AACTAMICON TOACACACA	120	ATCTITICCAC AGACTITICT GITTITIAGGG ATGAGACTAT TCTCTGCTTC ATCAAGGAAA	
GCTGGATAAA TTGACCGGTG TGTTTGCTCC TCGTCCATCC ACCGGTCCCC ACAAGTTGAG	99	5 AGAAAATCAC ATCCTAACAA AGAAGTCTGT CTAAGACAGT ACATCTCCTG TTGAACTTGC	55
MEDITIFICATION CONTINUES OF STREET TRANSPORTED STREET		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:	
TCTCCAGCC CAATTYCTAC GCGCACCGGA AGAGGGAGGT CCTCTTTCCT TGCCTAAGGC			•
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:	50	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	50
(D) TOPOLOGY: Linear		(A) LENGTH: 1285 base pairs	
		(i) SEQUENCE CHARACTERISTICS:	
			ř
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1290 base pairs	45	(1) THENDEMENTAL FOR GEN ID ND - RO.	45
(2) INFORMATION FOR SEQ.ID NO: 81:			
	1168 40	aaaaaanna aaaaactcga ggggggcc	40
	1140	GITACATATG TATATCAGAA TGTAAGAAAA AAAAATTTAT TTAAAAATAT TTITGGCAAA	
aataogaaaa taataaacat tgaag	1080	ACCITITICC TITITITITI ITITITICIOA ITIGALITCIS GIFACAGIGC CALAAACCIT	•
AACTAITITI CIGIATICCA AGAGCIGAGA ICTIAGAITI TAIGIAGIAT TAAGIGAAAA	1020 35	CAMATTIOCC AGCITGGAGA INGAMAGAA ITCAMCARIA TAICAMARAC ITTCCTTCCC	35
ACATOCITIAA TICACAAATG CIAAITICAT TATAAAITGT TITIGCIAAAA TACACITIGA	096	THIGHIGITI TANGTOTICCA AACAAGITAG ACCTOCCAGC TGAATGATAG ACAAATAGTG	
AACTGACTGA TAAAGCTGTA CAAATAAGCA GTGTGCCTAA CAAGCAACAC AGTAATGTTG	30	TITITITICAA ATAGTGAGGA ACTGACCATT ATATGCCTTC ACTGGCTTCT TGTGCAATAA	30
AACTIGCCCT TCATTACATG TTTCAAAGTG GTGTGGTGGG CCAAAATATT GAAATGATGG	840	GAITTIGIGI GITICANGCI CIGGAITITI ITTITITIC CITCICIOGG ITTIAAGAGAI	
TICCAAAAA CAIGIACIGA CIICCCGIIG AGIAAIGCCA AGIIGIIITII TIIAITAIAA	780	gocaccttaa Gtostictaag Raticcticag gcatictita aggaraaaa ggataccttt	J
CAINAACAGA AITIAGGIAGI ATATTGAAGA CAGCATCATT ANACAGITAT GITGITCTCC	720 25	aaacaaaa cccaaccaa ccaaccctg ttgtgctcac tgstgcaaag agaagatcag	25
ACTROCESSES CAAAGAAACTGAC CATAAAAAA AATTACCTOS TOAGAAGTTO	099	ACTAMATHGA AMGACATTG AATCACCAAG GCCTGGGAITC AACCTGGGGT GTCCACAGA	
GINSSITITA TICGACATGA AGGAAAITIC CAGATAACAA CACTAACAAA CTCTCCCTTG	600 20	TCTGTGCAGG TTCATAGACC GAAGATACTA CACACTTTAA ACCAATTAAA AAGAACCAAA	20
GGACCTITIGI CITCICIGISA AATGGIACIA GAGAAAACAC CIAIATIAIG AGICAAICIA	540	ATACAACTIG TITICICCCCC TITICCTITI AAGCTATIIG TAGAGITIAT GACTAANAG	
CTIVITGCTC TITIGGIATOT APAICISTICA ANGITITAAT GAICTGCCAT AATGICTTGG	480	Caaacigaca aattiaaggag ottaaagaag taatttitti aagccaacaa taaaaatata	-
GOOTOTOTOT OTOCOCCAT ATGININING ACAGGCACAT CITITITACT ITTICIAAAAG	420	ITTATITITIA ACAAGIGIAA GAAGACIAIA ACTITIGAIG CCAITGAGAI ICACCICCCA	7
ATCCAGAAAC CAGTATCTCT NAAAAACAAC CTCTCATACC TTGTGGACCT AATTTTGTGT	360	TICHTITIT TITIGIAGII GGGGGGAGT TIGIGAATGG AACAAACTT GTITAAACAC	
ANGTICCICG ATCTAGITICA TATTTAAACA ATATCTAGIT GATATTITCTC ATTCAGITICG	300 10	GGATTGCTGC TCTCCATTAG GAGACAATGA GGAAGGAGGA TGGATTCTGG TTTTTTTTTT	2
AAITICITATISC TOCCTONGCC CTAGAAITTT CCCACCAACG AAITIATTCCA GSTAGATCCT		ATCCTTACAC CAAACCTGAT GGCGTGGAG ACGGTTGTGT GACAATGGTC TGGATGGAAA	
GACTCCTCAG ACTICACCTA ACTITIGGAA CICTCTITITG GAGGCTTCTC ATTITCCCCCT	180	AACTICATGG GGCAGTTGCT AGAGTTCGAG GAAGACCTAA ACAAGGGTGT GACACCGAGA	
OCCUPACT GRAFFINGT GOTIGCACTT TCATFIGCTT TAGFICTAGA ATCACCTGTT	120 5		
CITCHITCCC TITCCIONCE CAGGARICAG CIGAGARITC AFFICGATIOF CATGCCTCIA	09		
CICCAGGACA GITTGGTCAG AGCTGCAAIT CITAGTCCAT GGTCTAATGC FICAGTATCT	**		

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CATTCAGATT GATTTAGAGA CTGGCAAGAT TACTGATTTC ATCAAGTTCC ATTCACCCAG GOTGACTCAT GATOCCCGCA CCATCCGCTA CCCCGATCCC CTCATCAAGG TGAATGATAC сттемансее сининате наноситось возсовоють вазовнего анситивае GGAGCAGAGC CTGCCGTGCA CCGAGAGGAA GCCAGCTGCT ACTGCCAGGC TGAGCCGTCG GCAGCTACTG CTTTTCCTCT GTGCCACCCA CTTTGGGGAG CCATTAGAAA AGGTGGCCTC CCAGGTOOTC TOGTCACCTC AGAGGCTCCG CAGACTCCTG CCCAGGCCAG GACTGAGGCA CAAGTACAAG TIGTOCAAAG TGAGAAAGAT CITTGTGGGC ACAAAAGGAA TCCCTCATCT COCCCCCAC AGCCGCCAGA TCCCCGCACC CCAGGGCGCG GTGCTGGTGC AGCGGGAGAA aggaacerea erarecese ecceegadas erecegadase ecceasease cagacerare AGCCTCAAGG CACTICIAGG ACCIGCCICT ICICACCAAG,AIGAACICAC IGGITICIIG TITATIGIAT ICIGIAACIA TAGAACIICI ATTIWATICI TITTIGGACI IGCIAAGIIG AAAAAAAAA AAAAAAAAANC TCGGGGGGGG TRAGGGRAGGG GGCGCTGGRG CTTCCRACCC GRGGCRATRA ARGARITOTT GCGTRACTCR ACCAGGGAAC CACGGCAGAA GCGCTGGGGCG GGGCTGAGGG CGCAGGTGCG GEACCTGCCG AACTACAACT GEAACTCCTT CEGCCTGCGC TICGGCAAGC TOTOGOGAAT TCTAGACCCA CAGGCCAGCA GCTAGAATCC CTGGGCCTCC TOTTIMATEG TITTIMAGITO CATEGICAAG TITTICAGIAT TEACITATICE COTTEGACAT TIGIYCACIG TITICIGIIGA TICIWACICA IGGIATITIA AFICIICGIT WITITITITIC ATTACTCACC AGAGAAGATT TTTTTTGTTYT ACCARGTOCC TARGAATGCT AACAGTCTOG TOTTWADAWA CATTOTTIGA AAAATAATTI GGAGGAATAT TIGATTOTTA TGAACAAGOO GAGTTOTTTT ATAGACTOTR ATGATTCAAA AATOTTACAT OTTTTOGTAG TOTOTTTOAT INFORMATION FOR SEQ ID NO: 82: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (A) LENGTH: 684 base pairs
(B) TYPE: nucleic acid
(C) STRANDECNESS: double
(D) TOPOLOGY: linear GGGCAGTGAA TOGCCCCSOG 1020 1290 1260 1200 1080 960 900 840 720 660 600 540 480 240 180 120 360 300 69 35 30 25 20 5 . 70 3 8 8 8 SS TITITCITCC GAAAGATICC CCCAACAITA CCATTCCCCA CCIICCGIIG AATTITITIG TTATCAGAGC CCAACTICGA GGGCTCIGGG CITTAGCIAC TGICACCCCA TCATAACTGA COATGATGAT GAAGATGATG TICATGICAC TATAGGAGAC ATTAAAAACGG GAGCACCACA ecoccricas ecrecades regganases errogaeses ecrerimente eccecentra TECTIONALITY GRAGICTEGA GETTTETTEG TICOTTEGYC GOCGOOTTEG COCCETTETO GCTCTCATTT TGAATTTTTC AAGA GCTTCATGGA TIGATICICT TITTAICTIT CAGATTIICT TITAAAAAIC TIIGITIITT CTGCAGGAAT TCGGCACAGC TGCGCTGGAG GCTTCATCTT TGCCGCCGCT GCCGTCGCCT (2) INFORMATION FOR SEQ ID NO: 83: COGCOGGACC OGAGOGGATG AGGAGGAAGA GTOGCTCTAT GOCGATGAAA ATGAAGTTGA TGAAAAACAA AAGAGGATAC GAATGGGACT TGAAGTTATA CCAGTAACCT CTACTACAAA TOCTGATOTT TOTGATTATT TTAATTATOG GTTTAATGAA GATACOTGGA AAGOTTACTG AGTICCACIC TIAGAGOTAG ATTIGGATIC TITIGAAGAT AAACCATGGC GTAAACCIGG TOGAACTACA GOGACAAAAG TCAAAGGAGT AGACCTTGAT GCACCTGGAA GCATTAATGG GTATOGGAGT TATOGTACAG CACCTOTAAA TCTTAACATC AAGACAGGG GAAGAGTTTA TGAAAATGOT GTACCAAAAC CGAAAGTGAC TGAGACCGAA GATGATAGTG ATAGTGACAG AMBECCAGAA GAAGAAAATG CCAGTECTAA TCCTCCATCT GGAATTGAAG ATGAAACTOC AGTIOCOCIC GOOOCGACCA IGICOGCCGG CGAOGICGAG COCCIAGIOI COGAGCIGAG CAGGCGACGG GCAAATGAGA ACAGCAACAT ACAGGTCCTT TCTGAAAGAT CTGCTACTGA AGTAGACAAC AATTITAGCA AACCACCTCC GTTTTTCCCT CCAGGAGCTC CTCCCACTCA Ê <u>£</u> SEQUENCE CHARACTERISTICS: GTACAGCAGG GAAGAACTGG AAACTCAGAG AAAGAAACTG CCCTTCCATC GAGITTACIT CICCICCITC SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 2024 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AUGITAICGG TCAGACTATA ACTATCAGCC GAGTAGAAGG TITIGITICAAG ACTOGGCTTC CACCGAGCAG

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960 900 840 780 720 660 600 540 480 420 360 300 240 180 120

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COGTOTGATO TATGACACCA AGGGTOGOTT TGOTGTACAT OGTATTACAC CIGAGGAGGO

420 360

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CTITICICCC AACCCIGIGG GAATARICAT YCATAICCIA RCTGCAGGCI ARAAGGIGGI GAMCACATAG AMCACCAGGT GATGAGACAA TCCTGGGART CCTGTTTTAC TTTGGSCCAT

660 600 540 480 420

TATAACCTAC CCTGCTGGAT TCATGGATGT CATCAGCATT GACAAGACGG GAGAGAATTT

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3	WO 98/39448 PC	PCI/US98/04493	*	WO 98/39448 PCT	PCT/US98/04493	_
:	305			306		
				GCCICATIGE TAAGTGIACT GCIGCCGAAG TICCCCCAAT ICCATGGGGT TCGIGTCTT	480	
	CETTOCACET CETCCATTIC TICCACETICE TECGACTOTIC AGACTICITE CACCITETAT	1140		GCCATCAACA AATACTGAGG GATGGGTTTT GGGACAGCTC CATGGGCATG GGGAAGGCAC	540	
v	TECACCACCO GOTTITECTE CTCCACCAGG CGCTCCACCT CCATCTCTIA TACCAACIAT	1200	v	TGAARCAGAG GACTATAAAA CATCCTTCTC TTATTCTCCA TACTGTCTTC TACACCTTTA	009	
ר	AGAMAGIGGA CATTCCTCTG OTTATGATAG TCOTTCTGCA COTGCATTTC CATATGGCAA	1260		AAGCCTGAGA ACTATACAAC CTTTCCCAGA CTCCCAAGAA GAGAAGAT TGGCAAATGG	099	
	TOTTGCCTTT CCCCATCTTC CTGTTCTGC TCCTTCGTGG CCTAGTCTTG TGGACACCAG	1320		GOCTOCTIGGG COCAGTOCTIG CTRATIGGCAA GTTTCTTTGA ATCAGGAAGG CAGGTGAGGT	720	•
10	CAAGCAGTGO GACTATTATG CCAGAAGAGA GAAAGACGA GATAGAGAGA GAGACAGAGA	1380	01	AAGGCCAAA TCACTCTCCT CCATAGCAG AAGCCATTTG GSCAGCTCCT TTGGTGATTA	780	
	CAGAGAGGA GACCOTGATC GGGACAGAGA AAGAGAAGGC ACCAGAGAGA GAGAGAGGA	1440		CANCITICOA TANCITITIAC ACITIACCACC TINCCAGCICT GITTINGCIGI GIAFITITICI	840	
2	OCOTICATCAC ACTICCTACAC CAACTIOTITT CAACAGGGAT GAAGAAGGAT ACAGATACAG	1500	15	TACARTAATT ITTITICAGCT ATACTIGCAG TITAATCAGG ATGGGTAGAG AGCTGTCCTC	006	
2	COANTATOCA GAAAGAGGTT ATCAGCGTCA CAGAGCAAGT CGAGAAAAAG AAGAAGGACA	1560		ATAAGSCTGG GGGTGGGAAG ATGGAATACT G	931	
	TAGAGAAAGA CGACACAGGG AGAAAGAGGA AACCAGACAT AAGTCTTCTC GAAGTAATAG	1620				
20	THGACCTCCC CATCAAAGTG AAGAAGGAGA TAGTCACAGG AGACACAAAC ACAAAAAATC	1680	20			
	THAMAGAAGC AAAGANGGAA AAGAAGCOGG CAGTGAGCCT GCCCTGAAC' AGGAGAGCAC	1740		(2) INFORMATION FOR SEQ ID NO: 85:		
ć	CHARGUACA CCTCCAGNAT AGGCATGGTT TTGGCCTTTT GTGTATATTA GTACCAGNG	1800	25	(1) SEQUENCE CHARACTERISTICS: (A) LENOTH: 825 base pairs		
53	TAGATACTAF AAATCTIGIT ATTITITCIGG ATAATGITTA AGAAATTTAC CTTAAARCTT	1860				
	STICTULE TRADIALGIA AAGTIAACTT TTTTTCCAAA ATAAAAGAGT GAATTTTTCA	1920		(D) TOPOLOGY: Linear	····	
30	TOTTARGETA AAAMCTETG TOTTOTACTA TITCAAAAF AAAAGACAG CAATGACTET	1980	30	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 85:		
	atatocaaaa aaaaaaaaa aaaaaaaaa aaaaaaggo ggoc	2024		COGGOCCOGC GGOCTOTICA GGTJACGGO CTGGTJACAG CAGCTCTACC CCTCACGAGG	09	
			36	CAAACHTOGC AGGGCAGAAG GACCAGCAGA AAGATGCCGA GGCGGAAGGG CTGAGCGGCA	120	
35			Ç	COACCCTOCT GCCGAAGCTG ATTCCCTCCG GTGCAGGCCG GGAGTGGCTG GAGCGGCGCC	180	
	(2) INFORMATION FOR SEQ ID NO: 84:			ACCICANCEAT COBSCOCTIOS ASCACETTOS TOSACCADOS GEOCTTETES, ESCECEGICA	240	
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 911 base pairs		40	ACCTOGRAGA GCTGTGCCAG COCCTCGTAC GCAACGTGGA GTACTACCAG AGCAACTATG	300	
	(B) TYPE: nucleic acid , (C) STRANDERNESS: double			TOTTOGRATT CCTGGGCCTC ATCCTGTACT GTGTGGTGAC GTCCCCTATG TTGCTGGTGG	360	
	(D) TOPOLOGY: linear		74	CTCTOSCIGI CITITICOSC GCCIGITACA TICICIAICI GCGCACCITIG GAGICCAAGC	420	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:		.	TIGHECIETT TECCERGAG GTGAGECEAG COCATEAGTA TECTETECT GGAGGEATET	480	
	CECECCHATA GCCGGACGGG GATCTGAGCT GGCAGGATGA ATGTGGGGGGT GGCACACACAC	09		CCTTCCCCTT CTTCTGGCTG GCTGGGCGG GCTCGGCCGT CTTCTGGGTG CTGGGAGCCA	540	
S	GAAGTAAACC CCAACACCCG AGTGATGAAT AGCCGAAGGA TCTGGCTGGC CTACATCATC	120	20	CCCTROTIBET CATOGRETIC CACRETIBECT TECACCAGAT TGAGGETGTS GACGGGGAGG	009	
3	THOOTAGAAT TOCTOCATAT GOTTCTACTC AGGATCCCCT TCTTCAGCAT TCCTGTTGTC	180		ACTICAGAT GAACCCOTG TCAGGTGTCT TCTGGGACCT GCCGGCCTCC CGGGCCAGCT	099	
	TOGACCETTSA CCARGOTCAT CCATAACCTS GCTAGGTATG TCTTCCTTCA TAGGGTGAAA	240		accessocc reconnece reportecte escretecte crossocca, escocione	720	
55	GEGRACICET TTGAGACTCE TCACCAAGGA AAGGCTCGGC TACTGACACA CTGCGAGCAA	300	SS	CCATCACAAG CCCGGGGAGG GATCCCGCCT TTGAAAATAA AGCTGTTATG GGTGTCATTC	780	
	ATGGACTATG GGCTCCAGTT TACCTCTTCC CGCAAGTTCC TCAGCATCTC TCCTATIGTG	360		AGGIANAANA AAAAAAAAGG GGGCCCCTC TAGGGGTCAA AGTTA	825	
9	CTCTATCTCC TOGCCHGCTT CTATACCAAG TATGATGCTG CGCACTTCCT CATCAACACA	420	99			

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(2) INFORMATION FOR SEQ ID NO: 87:

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(1) SEQUENCE CHARACTERISTICS:

TTCACTTOCA GCCOTGATGG AMAOTINGCA TGGAAGCTGA GACTCTCACT GACAGTGAAA AGGTTOTTOC CCAAAACACT GAAAAAAAACT GCCCTGGCCC TGAACCAAAT ACCTTGAACC CTOGCCAGTT GCATTTCCCC TGCAGGCTTG AGCCCAAGCC AGAGCCTTGA AAAGGTATTC TOCCCACITY TYTTCTOTOT CCTGACAAAG AAACACAGAG TAACTTGATT GCCCTGTGAC COCTOMANTO AMEMICANTICO CTGCTTTCCT GCCAMGGATC CTTGTAGGGT NCCCCCAGCT TOCTTIGGAC TGATCACCCT GCCAGTCTTT TGTCTTGGGC AATCTATACT TTTNCTCAGA CTCACTOTCT GTTGTGAGGA TACGCTGTAG CCCACTCATT AAGTACATTC TCCTAATAAA CTCGTAAACT CCATACCCTG ACCCCCTTGT TTTGGATATA CCCAGGTAGA ACAACTCTCT OCCUTTAGET TEMAGAATAC AGGATEACET GTACCEMAGE CETTAGETEA AGETETGETT TTACCTTATG CCCTCACTTC CTGAGTTAAC CTCCCAAATA CAGGATTCAC CTGTACCCAA GOTTECCAAG GECTACTGAA GOGACTTAAC ATACTETTAA TEGETTTECT CTCTETTETT TOGRAGARCE CARACTARGA CAGTGCTCCT GGTGCCCT 1020 1200 1080 720 960 900 840 780

> 1140 1080

1020

960 900 840 780 720 660 600

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45 6 ઝ છ 25 8 5 5 8 CTOCTCCTCC TGACCCGGGG CCTGGTGGGG GTCGGGGAGG CCAGTTAFTC CACCATCGCG TIGCCACCCC TGAACCCCAC CICGIGGIGG GCAGAICIGA GGGCICIGGC AAGAAAICCI TACTITICCCA TICCGOIGGG CAGIGGICIG GGCTACATIG CAGGCTCCAA AGIGAAGGAT CCCACTOTCA TIGCOGACOT CITIGIOGCO GACCAGOGG ACCGGAIGOT CAGCATOTIC ATTOCCTICT GRICCCTOGT GACACTOGGG TCATCCTTCA TCCCCGGAGA GCATTICTGG GAGACCCTCC TOTCCATGAA CTOGGCCATC GTOGCCGACA TYCTGCTGTA CGTOGTGATC CIGICCCITG CCIGCGCCC IGGIAGCAIC GIGGCCACIT ANATITICAT CITCATIGGA connected appropriest chalectary descripting deficienter culteriories CTOTOGOCTC COGCATTCCT OCTOCOTTCC COCOTOGTCC TTOGOGAGAC CCCACCCTOC ATGGCTGGAG ACTGGCACTG GGCTCTGAGG GTGACACCGG GTCTAGGAGT GGTGGCCGTT TOCTTOTION COGNOTICOG GGCTCTGCAG TICTCGCTCA TGCTCTGCGC GTTTGTTGGG GOTGGGAGCC COTACCTICAT TOGCCTGATC TOTGACCGCC TGGGCCGGAA CTGGCCCCC CCTACCCGAC GCTCCACCGC CGAGGCCTTC CAGATCGTGC TGTCCCACCT GCTGGGTGAT CTICCCOGAG ACTOCIGCIC TICCTOTGAC AGICTOATCT TIGGACICAT CACCIGCOIG CTOCTOCTOT TOCTOGTAGT GCGGGAGCCG CCAAGGGGAG CCGTGGAGCG CCACTCAGAT GGCCTAAACC ACAGCTOCAC ACCOGAGICC TODGIGIOGOG CCITOGGIGIO GAGATCAGCC OCCGOCICCG CCACICCAAC AGITICOICC IGICIICCCI GOGCTICACI GCIGIGOCCI TIGICACGO CICCCIOSCI CAGACACTAC ATGGGTAGCT CAGGGGAGGA GGTGGGGGGTC CAGGAGGGGG ATCCCTCTCC GCGCACTTCC TGGGCACCGC CATCTTCATT GAGGCCGACC GCCGGCGGC CCTTGGCCTG GCCCAGCTTC CAGAGGGACC CTGGGCCGTG TGCCAGCTCC ACCIDEACAT CIGCEACAGE IGGEECIGGG COCACCECAE GAAGGGECTIG GOCCOCTICEA COCGOGIGEC COTGGCCAGT GTGCTCATCT GGAGAGGCTG GTGCAGGGCC TGCTGCACGA AGCAGGGTCC ACAGACGACC GGATTGTGGT

30

GGGATGTCCT TTGATGGCAT CAAGACTTTA GCTTCTGGTG CGCTGTGTCC CAGCTCTGAT

TETTETETEA CITITETECT TITIGECGATT AGTOGACGIG ACAGAGATOT GAATGOGGCA

25

GCTGGCTCAC TGCTCTGGCT TCATTTTCCA GAGCTGCCTG CTGCAGTCAC ACTTAGGTCA GICTOGGACA CTOGCCTICC TOCTTACCTO CTCTTTCCTT CCTCCTTGGT CGGAGGAGG 20

CAGGAAGGGT GTAAGGAGAG GATGGATCCT GATACATGGA TTCAGGATCA TTAGGGTCCT TOTGGGGAAG TOGGATOGCA GCATGGCAGG GCTTTTGGAAA ATGAGAGGTG AGAGTKTKTC CTOCCACCTA CTOGAGAAGC CATAAGCTGC AGCTTTAGGA AAAGGGAACC COGGGCAGAG

300

180

15

GOGACOGITT CTOGCTCTCA GGCTCTGAGA AGCTGCAGTT TATGAGTGGC TCTGTGTGTG CATGIAAAAG GAIGAAAIGI GACIICIGGI GIITITITIAI IICIAIGGAG GGACIITICIG 5

g 0 9

TOPOLOGY: linear STRANDEDNESS: double TYPE: nucleic acid LENGTH: 1238 base pairs

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86:

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INFORMATION FOR SEQ ID NO: 86: (1) SEQUENCE CHARACTERISTICS:

SEQUENCE DESCRIPTION: SEQ ID NO:

240

180 120 60

480 420 360 300 (A) LENGTH: 1460 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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AACAGGGGCA GCCCCAAGGG CTCGGTGCTA TTTGTAACGG GATTAAAATT TGTAGCCAG

1380 1320 1260 1200

1440

(1) INPORMATION POR SEQ ID NO. 88: (1) SEQUENCE CHARACTERISTICS: (1) LEAGUINE CHARACTERISTICS: (1) LEAGUINE CHARACTERISTICS: (1) LEAGUINE CHARACTERISTICS: (1) LEAGUINE CHARACTERISTICS: (1) LEAGUINE BESCHETTICH: SEQ ID NO. 88: (2) STRANDERSES: duable (2) STRANDERSES: duable (3) TOPICACY: Linear (44) SEQUENCE DESCRIPTION: SEQ ID NO. 88: CAGGRICOLAN GYOGRANOTO TRANCOCCIA ATCHTOGACT CHARACTERIC CHAR

330

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 89:

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(i) SEQUENCE CHARACTERISTICS:
(A) LENSTH: 1186 base pairs
(B) TYPE: nucleic acid
(C) STRANDENESS: double
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 89:

9 120 180 240 8 360 420 480 240 1140 9 9 720 780 840 900 960 1020 1080 1186 GOCACGAGOCO GGCAAGCOGA GCTAGGGTGA AAACTGGGGG CGCACCAGGA TGTNNGACAG AAAAGCAGAA GATGAGACTC TOTTCATTCA CTTTTCCTAG GCCCATCCTG TGGTCATCTT ICCCCTCCC ATCATACCTC CTCCTTCCTG GAGCCTCTGC CGGCTTGGCT GTAATGGTGG CACTTACCTO GATATTTCAG TGGGAGGATG AAAGGCGAGA CTCACCCTAC GCGGTGGGAC AGATGGGGAG AGGAAAAAGG CAGAGATGGC CAGGAGAGGG GTGCAGGACA AACCAGAGAG CITICGGTCAG GGGAAAAGGG TGGGGAGAAA GAGGGGTGCA GGCCCTGCAG GCCGGTTAGC CAGCAGCTGC GGCCTCCCCG GGCCCTTGGC ATCCAACTTC GCAGACAGGG TACCAGCCTC AGCOSTIGATO STOCACCATG ACATOCAACO OSTATATATA AAGATAAATA TATATATATA GIGGCTIGIC TTAIGTCGIG ATAGCACAAG IGCCAGICGG ATTGCTCTGT ATTACAGAAT CANICICITY CICTOTITICY CICCICCCCA COTOTICACCC TICCCCTOTIC CATOTICCOC CIGGIGIGIA ICATAGGAIT IGIICACATA GIGITAIGCA IGAICTICGT AAGGITAAGA TOTATICTANA TTATGGCACG AGAAATTATA GCACTGAGGG CCCTGCTGCC CTGCTGGACC AAGCAAAACT AAGCCTTTTG GTTTGGGTAT TATGTTTCGT TYTGTTATTT GTTTGTTTTT AGIGITITIA AITICAAT GIICIAGITA AIGICIACCI CAGCACCICC ICITAGCCIA ATTITAGGAG GITGCCCAAT ITTGITTICIT CAAITTITACT GGITACITIT ITGIACAAAT receseere ecercises rendacises esterearin engrecaene carreneren COCTICTICE TRACTICATE TRACCECTICE CCAGCICACT TOCOCGAGIT GTGCTTGCCG CICCITIATICI GITICIAGITIC CGAAGCAGIT ICACTCGAAG ITIGIGCAGIC CTGGITIGCAG CITICOGCAT CIGCCITCOT ITCGIGIAGA ITGACGCGIT ICITIGIAAT ITCAGIGITIT (A) LENGTH: 1821 base pairs (B) TYPE: nucleic acid (C) STRANDELNESS: double (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 90: 35 15 2 25 8 6 45 S 55 8

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SEQUENCE CHARACTERISTICS:

(A) LENOTH: 696 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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AAAAACTCGA GGGGGGCCCG GT

1380 1320 1260

2 INFORMATION FOR SEQ ID NO: 92:

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1680 1620 1560 1500 1440

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55 TACACATTIC ICATATIGGG GIICGACAGG TAAACACAAA CIGCIATTIC AGIAGAAAA

TCATOCTITIC CITICCTAAR TITITICACAF ACCITITICT ATCACACICE CITICCTITIC

ACAGATTAAA GATACAGTTA CGTAAACAGC AAAGTAATTT TATAGTGCTT CATCCATTTA

CATTOTICO TOCOTOCCAT CAGGICAAAT CAGGAGGOIG CAGIGAATGC CIGITOTITIG AATOTOTAGC AGTICTICCT GTAACTCTTT AAAACTTOGC TATAGGCTOT TIAGCACAGT

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AATTAGAGCA AAACAAATCC TCTACAAATC CAAGGCAGGA AAAGTGGTGG CAGAGTGACT TITICATICCC TATAAGCAGG TACCITAGTA GGGCAGATAT AGGAAAAACA

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ANGANTAAGT ATCATGTGAT TATTTTAGCT TTACAAAAAA AAAGTTGAAT CCATGATTCT CCTATAAGGT AACTCTTTAG TCCTCCATTT AGCACATTTT

1140

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TITTAAAATG TGACTCTCCC AGAGAAGAAG CCCCTGGCTG TATGAAACTT GACGGCCCTT

TIGINAGGIG CCACCCCAA ACTITANGGI

AGCTANACCA

ATTITITAAAA GATTCAATGG

CTCACCCTCA CAATITATITI CCICCICCCG IGCCAGCCCT ICITITIGIGI CIGAAACCGG GACGITITIC IGCICICITC IGGCCCICCA IGGAGCCAIG GGCCICGGCC ICGGCGGCTC AGCIGAGCIT ICITATICCA CCCTITCIGG IGICTATAGG AATGCATGAG AAGACCCIGG GCCAGGAGCT TICICAGAAA CAGICATAAA CGAICICITIG AGICICITIC TIGICCICCC

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420 360 300 240 180 120

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TOTTTCCTGT TITCTGTGCT TICCTTTTTA CAGGACTCCC GGAAGGCCAC TCATGGCCAT

TACCACCOCT TOGAGTCTCC CGAGGACACA AACAGGCAGA GAGGGACGTG TAGGGAGAGT AGCTIACICA COCCGGAGIC TITICITICI CIIGCICCAA GAAGAGCCCI GIIGGIGCII AAATGACTIG ACCTIGCCAT CIGIGITICAA GGICACGGIT IGCIGIGGGG TICCIGGGAG GGACTGGAGA GAGGTGAAGA ATTITIGCAGG TGGGAGATTT GGATTTGAAT GTGGACTTGT

1200

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TOGTOGTICCT GATTTATAGG ATTTICATAAT TAAAATGTICT GCTGAATAAA AAAAAAAAAAA CITGITICATC CICCAGATGT AGCTATIGAT GTACACITCG CAACGGAGIG TCIGAAATIG

862 840 780 720 660

6

TICCAGGIIG ATAATGATGG CCAGAAGATT TAACATACAA AGTAATTCTC AATGTAAAGC TATTCAGCTC AATGCCCTGT AACCCACCCT GACCTTCCAC ATCATCTTCA AAAAGCAGTT 1080 1020

ACAGTTTCAT ATGAGAAAAA TIAAGAATAA CTATAAAATT GTTAAAATAT CCAATAATGG 960

35

GCAGTGAGGA TGTTTCCAGA ITGTAGAGAA AATCCCCITT AGCTACTTAC CTACATOTGA ACTITITIGGA ACAGTAATGG CAGCTICTAG TACAGCCATT AIGIGCCATT TICTIFICCTI 840 900

TACTAAAACA AATGACACTC TAAGAAAGTT TGGGAGCCCC ATGCTGAGAA CCATTTCTGT

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780

TGACTCACCT TAAAGTICCT ATIGACATCI ACIGCTITTA AACCTATITG AAAACTCTGA

CTARTIAGAA GGGGAAGTIA GCCACAGAAA ATCAACTIAT CTATAATTAC AAAATTCICT 720

CAGCAGNAGO AACCCCACCA GCCTAAGTCC AGCAGAGGAC CTCCCACCCA ATGTCTTGTT 660

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TOCCCTTTTT CCCACCGATT CGGGGCNTGG TGAAGGTGGG AGATGTGAAC TCCAATTAAG

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 91:

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(2) INFORMATION FOR SEQ ID NO: 91:

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SEQUENCE CHARACTERISTICS:

LENGTH: 862 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double (D) TOPOLOGY: linear

AACCCGGGGG GGGTTTCCCC C

GAGGGOTTAT TITNOGCTTG GGCACTGGGC CCTTCGTTTT TACAACGTCG TGANGGGGG AAAAAAAAA AAAAAAAAAA AAAAAAATTC CTGCGGGCCG CANGCTTTTT CCCTTTGGGT

> 1800 1740

1821

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20 GROSCIATICS CTACAAGGCS CACSCAAGGG ATOCSSAAAG CSSAACSTTS TSSAGGGSTS 540 600

AGICAAAICA CTACTICCAI IOCTACTITA GAICAGCCAA AGIGGIGACI GCIGCAGIGI

15 CACCCAGACA CAAGCCCCTT TCCCAGGTCA AACCACAGGC CGATGCATCT CCAGTTTGAC 480 420 360

TGATTCAACA CAGCTOCCCA CACAAAGCCA GTOGTAATAC ATCTGTTTAC CTTTCCCTAT CICCATOCTA GAAGCCAGCC CTAGGMAGCT GCAGTTACTC CCTGTGACTC AGCAGCAGGC 300

GEAGITICAC ACACAATEAC AGGCTGCTOG GGGACATTGC AGGACCCCTT TTCCTYTCCT 240

5 KATCCOTCCT CACTOCGCTC AAGATGGCCT CAGCAGACAC CAGTTACCCA GCTGAAAGTC ACAATCCCTC CCAGAAGTCT CCCAACACTA GTGCTGACCA GAGGTGGGGC TCTCAGGCTA 180 120

SEQUENCE DESCRIPTION: SEQ ID NO: 90:

(D) TOPOLOGY: linear

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AAAACATOCT TICAGGGCGT CCCCTATGTA TICGGGGGGC CCAGGGACAC TCAGGCTGGA

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	A APPREASED PROPERTIES. ED. Th. ND. 92.			
	(X1) Seguine Description: On the Co.	;		CTGAAGTCTA TATCGGCATC GGGAAGCCTG CAGAAGCCAC AGCCTGTACC CAAGAAGCTG
	CTGAGGCGAG TGAAGTGGAC TCTGAGGGCT ACCGCTACCG CCACTGCTGC GGCAGGGGCG	9 .		CCAACTICIT CCCAATOTIC CACAATOTIC TCTACATGGG CGGCCAGATT GCTGAGCTCC
S	TOGAGGCCAG AGGCCGCGG AGGCCGCAGT TOCAAACATG GCTCAGAGCA GAGACGCCG	120		5 GOSGAAGCAT GOACGAGGCG CGGCGGTGGT ATGAAGAGGC CTTAGCCANT CAGCCCCACC
	AAACCCCTTC GCCGAGCCCA GCGAGCTTGA CAACCCCTTT CAGGACCCAG CTGTGATCCA	180		CARGIGAAGA GCATECAGGS ACTIGGGCCT GATCCTTCAC CAGTIAGGCC GYTACAGTYT
9	GCACCGACCC AGCCGCAGT ATGCCACGCT TGACGTCTAC AACCCTTTTG AGACCGGGA	240	01	_
⊇	OCCANCIANCE OCCURITIONEC CINCOLOCOCO TOCOCOLITIC OCTUCIACOCT CALCITOCOTO	300	•	
	CTTGCAGCCC TCGAGAAAGC TCAGCCCCAC AGAACCTAAG AACTATGGCT CATACAGGAC	360		GALACTITIC GACTICEAGE CTAGABATIC CONTENSION TITLACTAROA TECHNOLOGY
15	TCAGGCCTCA GCTGCAGGCAG CCACAGGCTGA GCTGCTGAAG AAACAGGAGG AGCTCAAGGG	420	15	
	GAAGCAGAG GAGTTGGACC GAAGGAGCGA GAGCTGCAGC ATGCTGCCCT GGGRGGCACA	480		ACCACACATE GRECONTICE CICAMAGESCA TRANSPICIOC GAACTEMOCO CAGGAAGAA
ç	GCTACTOGAC AGAGAATTIG GCCCCTCTA CCTTCTTTTT GTCCAGTTCA GCCCTGCTTT	240	20	_
₹	TICCAGGACA TCTCCATGGA GATCCCCCAA GAATITCAGA AGACTGTATC CACCATGTAC	009		
	TACCTCTGGA TGTGCAGCAC GSTGGNTCTT CTCCTGAAYT TCMTCGSCTG CCTGGGCAGT	099		
25	TCTGTGTGGA AACCAACAAT GGGGAGGCTT TGGGTT	969	25	
				TAITITIATO AGAGOTOTAA AGAAAGAAC TOTOCTITITO TOCOCAGCAC COCTOCTGCC
30	(2) INPORMATION FOR SEQ ID NO: 93:		30	CCACTITICAC CCAGAAACCA AATGIGAACT TCCTGTCTCC CACCTCAGCA CTAGTCCATG
	(i) SEQUENCE CHARACTERISTICS:			CCAGGACACC AGCTGACAAT TTCTTGGTTT TACTGTCAAT AATTGTACCA TGTGATCAAT
	(A) LENGTH: 1886 base pairs (B) TYPE: nucleic acid		36	TACTOTICCIC ACTINGAACA AGCCTGAGT CCGAGAATAT TTATATTITA CCAATATATO
35			C	CCTOTTACAA GAGAAGGAAA TATGAGITAT TIVAGITIAA CTTITITAIG IGAAITICAGA
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:			GITTAITIAL CCAGGGAAAT AIGTACAAAG AAGCTICAAA 1GGAATAITT ACCGACAITC
9	CAGGCCACTG ACCTITCTIT GCGAGGGATG CAGGAGGTCC TACAGAGAAA GGCGCTTCTT	09	40	CTTATACATG ACAGACACTT GOCTACATOG GAAGATGATG TTAATAATAA AATGATTTT
	GCATHICAGA GGGCCCACAG CCTGTCACCC ACAGATCACC AAGCAGCTTT CTACCTGGCT	120		aantogaaaa aaaaaaaa aaaáan
į	CICCAGCITIG CCATCTCCAG ACAGATCCCA GAGGCTCTGG GGTATGTCCC CCAAGCTCTT	180	45	
.	CAGCTICAAG GIGACGAIGC CAACTCCCTG CACCTCCTTG CCCTCCTGCT GTCAGCACAG	240		(2) INFORMATION FOR SEC 1D NO: 94:
	AAGCATTACC ATGACGCTCT GAACATCATC GACATGGCCC TGAGTGAATA CCCAGAAAAT	300		(i) SPOILFAUE CHARACTERISTICS:
20	TICATACTAC TOTITICCAA AOTGAAGTTG CAGTCACTCT GCCGAGGCCC GGACGARGCA	360	50	
	CIGCIGACIT GIAAGCACAT GCIGCAGATA TGGAAATOCT GCTACAACCT CACCAAGCC	420		
,	AGTGATTCTG GACGTGGGAG CAGCCTCTTA GATAGAACCA TTGCTGACAG ACGACAGCTT	480	55	(xi) · SPOUR
Ç	ANTACANTA CITTOCCAGA CITCAGOGAI COCGAGACAG GOTOCOTOCA TOCCACATOG	240		CTCAGCT
	GTAGCAGCCT CANGAGTIGA GCAGCACTG TCGGAAGTIGG CTTCGTCTCT GCAGAGCATG	009		CHOCHCHOC COOLEANNER AGAINMENT PPINICADAC TARADETAT INCOMPINICAT
9	CCCCTAAGCA GGGCCCGCTG CACCCTGGA TGACGCTGGC ACAGATCTGG CTCCATGCAG	099	09	

9 120

8 55 GTATAAAAAA CACAGTTCTT GAATATTAAT GAGGAGGAAC ATCTTTYCAT GTTTCTTGGC CATTTGCATT TCCTATTATG GAATATICIG TAATITATIT AATCCCCTAT GGATIGATAA TIAGGTICAT TATAGATAGA TTCATAACTG TATTTTCACC AAGTGTATGG AGAATGTTCA TTTCCCCCATA TAACCATACC ACATCCTATT CAGATGTCTT GAAAATGCCA TTATTTATTC CCTTAATTTT TTTCCTCTCG CTATTACATT GCCAAAGTAA CTCAAAAAAA GATAACAGGT CATAAAAACA CITTITACAT AAATAGGATC ICATAITICIG TAGCITITITA AAATTITIGGI AAGATTTCAC CAATTTACAA CTCCATCATT AGTAAGAATG CCTGTTTGCC TATAGTCTGC AATAACTAGA AATITATIGG ATCAGGITIC ACATITIGCAG TITIGAAAAC TACTACCAAA GOGGAACCIA TICCCIGIGG CITAGGIGAG CAIGIGACCA GOCCIGOCT CCIGAGICCC TTACIGIOGA TIGITIGIAT COCTTACCIG CITICIATIG GGITATGIGI GGATATATIG TGATTATAAA AAAAAATGGT GAGATTGGGG TTATTTTCAT GTTTATTGGC CATTTATAGT TACACTIGAT ACTITITATIC IGTIGGCCGA AAAAGAACCT TITCITATIT IGCATITICCC AAGGTGGATT TTTGGATCAA AGTGTAATTA ACATTCCTGT ACATGTATTT TGCTACTTGT GTGGGTATTT CTGTAGGATG GATICLAGITG TOCCTAAAGC CICCCITACT CCGGACTITA AAGITTTVIG AGCCAATAAA CCAAGGACCA TOTAAGCCTG AATTTGTGCC ATGTGGAGAG AGTCTGTCTG AGGAGAAACT GCCCTTAACA GAACTICIGC IGGAACTACT GGAAAGAAGG CIITAIGGAG AICCCAGGAA TITTIATING TICAGCANCI CCINCOCCAN CINCIGGIAA CACAACCITI ATTIATING ATATAAGITA ATAAAAITIAG CATGGCCTTC CATG GTATTTTCTT ATTIGATTTGT AGAAAACCTT TGTAATTTTA AATTCTAGAC TTTATGCACT GICCCITICI CCCATCCTAG CAGAAATCGA AAGAGAACTA ACTICICATC TCATITITCT CCACGCCCTA ACAGCTTCCT AGCCACAGTG ATAAAAGAAT GGGTATATAA CTTAAGCCAG GCTAAGGAAA ACAGITIAAC ATACATECTE TAGTCACCIT TECGTACGAA TATACATACA GCCCATTTTC CITTITITAA TTAIGAAAGT CTAATGACTA CCTTCTCATT TGITTAAGAT AATTGAATTG AGTTTCTGTT CTGATTAATA TAGGTTATTT CTITAAATTT CTTTAATGGT TGAATATGAT TAAATACTAT TGAATAGAGA GACCCTTTTC TCCAATGCTA CCAATCACAT TGTGCATGTG TGTGAATATT TCTTTAGTCT GGAGTCCAGT AGGGITTIGIT CICIGICCAC CITCAGICIT CCCAAAGGCC 1740 1560 1500 1440 1320 1260 1200 1140 1080 1020 1680 1620 960 900 840 780 720 660 9 480 420 360 90 240 177

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CCTCACCCTC TTTGCCCCCC AGCCTCTCCT CCTAGCCCAG TOCAACAKTG ATGAGAGAGC coroaccore corocactive econocacor ecocacectr economics ecitoconect CTOCCTOTOG ANTGAGGACA CCAGCACCCT ACAGTGTCAC CAGTTCCCTG AGCTGGAAGC TETETOGGAG GETGGCAACE TEACTGAEET GECCAACETG AGAATEGGET TETATAACTT GCTOTTCATO AGCATCATAG TCCTCGTGAT TOTGGTCATC TGCCTGATGT TATACGCTCT

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SCICIOSSEA GOGECEAGSE CITACICATE CICTISCITA TAGCEATOGE TOTOTICECT

720 660 600 540 480 420 360 300 240 180

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TOCTATOTOT OGNATGOOTC ARGOTOTOCC TOCCOGGOOCC TGGGTTTCTA GEAGIGGGET TECTOGETST KITCHTCTOTO CIGCIGGEAG GEGGEETIGG

GIGGCGSCIG

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SEQUENCE DESCRIPTION: SEQ ID NO: 95:

GOCACGAGEG AAGGCAAGGG GGCACCAGET CAGGACTGCA TETGCCTGCC ATTTCCCTTC

CACTOCTOCT TTOTOGRAGIC TGACATTAGA AAGOCAGCGA GAAGGAAGAT TCAAACAACG

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AACCCIGATT TOCTOCTICE CONTINUATE AGIGTICCIG TEGRICICIGC ACCICCITIC

TOTOCOCOG CAGAGGGCAG TAGAGATGGC CGGCCCAAGG CCTCRGTGGC GCGACCAGCT

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ACTOGRACTE CARCECCAGA GGTECAGGAG TGATETETGA GTGACTERAG GGACATOGIT AAGGAGCITC CACTCAGCCC ACCATAGIGA GIGGGCCGCC

ACAAAGACAA GGCTTGACTG CTTCAAAGCT TCCCTGGACC

ATGAAACCTA GCAAAGAACT TACGGCAACA

TGATTGCTGT CATGGGGCCA

GACTICCAGG CIGATITICC AAATGCCAAA AACGAGGACA TTAAAAGAGC GAGCACCTCA

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8

CTCCACTICC CAACCCAGAA CTIGGAAAGA CATTAGCACA ACTIACGCAT IGGGGAATIG

1560 1500 1440 1380 1320 1260 1200 1140 1080

CCTCGCTGTG AACAGGACTG GACGGTTTGCG CACAAACAAA CGCTGCCACC

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ACAMATCACT CCCATGATGA GACCCTGGAG

2

ACACCAATTC CIGCITTAAT TAATGGATCT GAGCAAATCT TCCTCTAGCT

TCAGGAGGGT

6

GICTGICCIT

CARAGAGGCC GAGGGGCAGC AAGGGCAGMC AGGGCACCTG TGACTTCTTA GTACAAGATT

CAGGACTICC AAGGCTCCCA AAGACTCCCT AAACCATGCA GCTCATTGTC

1020

960 900

TAACGCTGAT CICCAGCTCC AGCGATGGAA CCCACTACAG AGGAGGTGGG GCCCCTGTGT GOGTTCAGTT CCAACCATGG TCAGAGGTGG CACATCTGCT CAGCCATCTC ATTITACAGC CTGAGGGCTG AGAGGGCTGA GAGCAAGCTT GAGAGCTGCT AAAGGCTTAC GTGATTGCAA

840

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INFORMATION FOR SEQ ID NO: 95:

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SEQUENCE CHARACTERISTICS

(B) TYPE: nucleic acid (A) LENGTH: 2503 base pairs STRANDEDNESS: double TOPOLOGY: linear

	WO 98/39448 PC	PCT/US98/04493 W	WO 98/39448
	317		318
			GENCANCICC GODGCOUNGES GOCCUNTINGS INSTITUCIONES CAGOCOCCIAS CRISCOCCIAS
	TGIGTATITIT CTAGCACTIG TGTATTGGAA AACCIGTAIG GCAGIGAITT ATICATATAT	1620	GEOCCOGGO GETECCHARG CAAGGAGG CCCTGAGAGC TCCACCTAGT TCACAGGATA
•	TCCTGTCCAA AGCCACACTG AAAACAGAGG CAGAGACATG TACTCTGGTG TGATCTCTTG	1680 5	AAATCCCACA GCAGAACTCG GAOTCAGCAA TGGCTAAGCC CCAGGTGGTT GTAGCTCCTG
•	TECTEMBRICA CTETATERISES CTCCTGTECC TETAGETRA TAGCTAGETIS CECGGGGAACE	1740	TATTLATTE CONTINUE CONTINUE CONTINUE CONTINUES
	AAGCTACAGG TGAAAGCAAG GTAGCAGCTT GCGGGAGGAG GCCTGTCTGG CTTACCAGTC	1800	COMPACADA STOCKMENTED CANCESTIVENCY ACCANTIANCY PACTICIPANCY CARPANESTY
01	TATACACTOT GGCCTCAACC TCCCAGACAG GGCAGAGAAC TGTGGGCAGC TCGTTTGCTT	1860 10	GIINCHIMIN ON MUNICIPA DISCUSSION CHICARA DANIA CHICARA DANIA CHICARA MINISTERIA
	TCTAGGCTGG CTGGAGGGT GGGAGCTCAT TGATAGACTC ATGATGGAAA CTATTTTTGA	1920	AUGHITTI UMILAILI MAGMUANC LIOCANIII LUMAKIUMA MITUMAKAI managana cama sanam manambasa asanaman managasasa atamahan
-	AACAGGCTTC CTCCTTCAGG AGAGATCATG CGGACTAAAC TGTAGCAATT CCAGTGCACC	1980	TICHMINIC CITAVITACI TATATINA THEORY MINOCIPLE COLORAD
3	TESCHOTEAT CETTITETIT GEAAAGTACT GTETETTIGG TTECAGTAAG TIGGACCACE	2040	TCATCIATCA ACAGACCACA TCTATCCCAA ATTICICTA TATGGGAGGT GACGGGGAA
	ACATGACATY ATTITICCCTG GAACCTGGTC ACTGACTAAC ACAGACAATT GGGACTCCAG	2100	THE PROPERTY OF THE PROPERTY AND THE PROPERTY OF THE PROPERTY
20	AGCCTCAAGA GCCAGAGAGG GGCACAGTAC ATACAGAGGG AGTCAAATGG GATCTCATTT	2160 20	TICABAGAIG TOGGACIGAA TATGAAGIIA AAGAILAAGC TOCAAAAGGG GATGAAGIIA
	TEAGTECTICC CTTCCCCACA CTCAGAACGG CANCCCCAAG GCCCGGAGTG TCCAGGGCTT	2220	CICGRARACE STITCATOCA TITOTACICE TACTOGRAM ACTITISTET MACCINGRAM.
Ċ	CTEGCCTICAG GTGAATCTGC CAGGCCCAAG AAGGCACAAA GGTAGGAGCA CAGAGAGCCC	2280	ורשמפתישה שייו משמשונים שייו ביינים מייוים מ
3	CATTCCCACA GOCGGKOGGC CCAGCAGCAC CAGTGGAAGC TCAGCTGTCC TCCAGCTGCT	2340	TOCTGARISC CCIVITIUM ARICCIATIOS ATGACARITI ARTITIONICA GIAMARTING
	CTCGGCAGAC ACTTCAGTGC ACAGTITIATG CCCTAGCTGA AAAAGATCTC CCGGACGTAT	2400	TARAGITSAC AGGATCAGIT TIGGARAGATO CITOGARAGAR ARABIGARARO ALGARTATOG
30	TICAGGACAT CCTCTTCCTC CTCCTCCTCA GGCCTCCTGC TACAGGCAGA GCTGGAACCC	2460 30	aagaaattat tengagaatt gaaaggitu teetagatse aaaetuseagi agagatgiaa
	CCCGGCCTCT GGGNAGGCT GAGGCCTGGA GYCAGTGCCT GTC	2503	ACAGAITOCT CITGAAGCIT GIAISAACICC GGICAAGTAA CITGGGGAAG GICCAITGCAA
!			
32		35	CATITIATAC ATCTGATGGT GTTCCTTTCA CTGCAGCTGA TCCAGATTAC CAAGAGAAT
	(2) INFORMATION FOR SEQ ID NO: 96:		accaganti acttgaaaga gaggactiti ticcagatta tgaagaaat ggaacagatt
9	(1) SEGUENCE CHARACTERISTICS:	40	TATCCOGGC TGGTGATCCA TACTTGGATG ATATTGATGA TGAGATGGAC CCAGAGATAG
₽	(B) Tarvaint sout base passes (B) Tarvaint sout base passes		aagaagctta tgaaaagttt tgttttggaat cagagggtaa gggaaaacag taaagttaaa
	(C) STANDELNESS: COUNTE (D) TOPOLOGY: linear	-	TITCAGCATA TCAGITTIAT AAAGCAGITT AGGIATGGTG AITTAGCAGA ACACAAGAGA
45	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 96:	45	GCAAGAAAT GTGTCACATC TATACCAAAT TRAGGATGTT GAGTTATGTT ACTAATGTAT
	CTICGAAAGCC GAGGGTAGCC GAGCGGGGG GGCGCTCTGG AGCGGCGGT GCTCGGGCTG	. 09	GCAACTTIAA TITIGITTIAA CACTATCIGC CAAAATAAAC TITAITCCCT ATAACTIAAA
Ş	CEGIECGETE COCCAGNAGE ACCAACAGE CGAGCCGGG CCCGCCCCC TCCTCCTCA	120	atgigiatat atatataata gittaitaig tacagitaat ictactigitt tggcigcaat
3	TEAGGCCCGA GTGAGGGGG GCGGCTATAG CCGACCCGGCG GCGCCTTCCC CCCGCGTCCT	180	AAAATCGATT TIGAAATAAA TGAAATGTTG AAAATTTTGC TAGTTGGTTA GATGCTTATC
	ATCECCAACCO CACCACAAGC GOCCCCTOCA GOAGGAGGCC GAGGAGGAGG AGCATGTOGG	240	CITTAAATIC TACTITICIT GAGGGGAAAA AGICITICGIC TGGAAATACA TATTACTGCA
55	ACGSTITICGA TCGGGCCCCA GSTGCTGGTC GGGGCCGGAR CCGGGGCCTG GGCCGCGGAG	300 55	AAAATGTAGC ATCCTTTTTT AGGTAGGAGT ATTATAGCTT YCATTTTAGT TKGACATTTA
	GGGGCGGGCC TRAGGGCGGC GGTTTYCCGA AMGGARGGGR GCCTGCTGAG CGGRCGGGGC	360	GTGTCCCAAT GAATTGAATT TCAAATATGA ATCATAATCT TGAAAATCTT TAGCACTAAA
09	ACCAGCOGCO GCAACOCAAA GCCCCGGGCT TYCTGCARCC ANCGCCGCTG CGCCARCCCA	420 60	GICTIGGGAA TATATCAACA ACIGATITAC ATATGCAGAT GCTATITIGNA TACCAAGGGC

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AGGGTTCCCT TOGATCAGAC TCCTCTTTTT TATCCATGGC AGGACTGGGC ACTTGGAGTC acroandria recerereer reressoere enormande regrentiar escreteere CAGAAGGTTA AAGAGTOGTC TCTCATGATC ATGAAGACTT TGATAGTTGC GGTGCTGTTG GODACTICCA AAATCCATGA GCTCTACACA GCTGCTTGTG GTCTCTATGT TTGCTGGCTA AGCCTCATCT GCCTTACTTY ACCAGTATTT GCTGGCCGTT GGTTAATGTC GTTTTGGACG AATTYTOCAC TCAGGATATT TCTGTTGATT GTCTTCATGT GTATAACATT ACTGATTGCC CACCAAGCCA TACTCCAGCA GOGAGGGCCT GITTGGYTTTTC AGCYTTACCG CCGACCTTTA AATCAGCATG CICGAAATAA CAACGCTATT CCTGTGGTGG GAGAAGGCCT TCATGCAGCC TCTTATTTAT TGGGAGACCA GGAAGAAAAT GAAAACAGTG CAAATCAACA AGTTAACAAT TESCITUAAGS CECTEGTECS ASCOTEGACT GTGACCECCG GATACTTECT GGATCTTCAT GAGCTGCTTC TGCTTCAGGT TGTCTTGCCA GCATTACTCG AACAGGGACA CACGAGGCAG AATTITICITIC CATACAATOT CATGCTCTAC AGTGATGCTC CAGTGAGTGA ACTGTCCCTC GIAAATAGIT CIGITAATAA CCCACIGITT TACAITIGGT ACAICIGIGI CIGCTAATAC ACCATAAGGG CIGIGACGGT GAIGGIOGCA IOGAIGCCIC AGGGACOCAG AGIGAICTIC ATGGAGCCAA AGACAATCAC TGATOCTTTG OCTTCTAGTA TAATTAAGAG TGTGCTGCCT ATAAAGCTTA AAAAAAAAAA AAAAAAAAAA AAAAACTCGA G AGTIAGCITT CICACITITC IGCITGITTG TICAGICIGA ATTAAAATTA GACTITIGAAA TCAAATCTTT TICCAGCIAA CIAAAAACTG IGIACAAAAG GATIOCTIGI AAATAIGCAT TATTISTICAT ICAAGITITIC AICIGCITTA TAATIGATAC ACCITGAGGG ICACITTICI AACTGRCTAA CTICATTACC TIAAAGCCIA GAACATIAIT CIGCITIAIT TATAIGGCIT AATACTTITA CTATAATGIG GTACCACCIC AGCCCTAATA AATAATAITT TTACCTAATG TCTCACTITT ATTITIGTAGC AKGGGTTGCA TCGACTTTTT TACTAGAGAA TTITIACTAGA GACTICACIG GAAGATITAN ICCAANICHA GGAANIGHIC TITITHAITH TIATHYITHC TTTTTAAATG TCATGGGGGG GAAAAAACCCCA ACTTGGTGGA ACTCCCAGCT AAACAACCAA INFORMATION FOR SEQ ID NO: 97: Ε (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (A) LENGTH: 1631 base pair
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear LENGTH: 1631 base pairs 2801 2760 2700 2640 2520 2460 2400 2340 840 780 720 660 600 540 480 420 360 800 240 180 120 8 55 8 ટ 8 35 မ 25 8 5 5 S TCTTCCAAAA AAAAAAAAAAA AAAA GRAGCITICIO TICRATICCCA GCGGICCITA CCAGRARRAG CCIGIGCATG RARRARARAGA CATGCTCATC GTATCTOTGT TOGCACTGAT ACCAGAAAACC ACAACATTGA CAGTTGGTGG GAAAGSCCAC GTGAAGATGC TGCGGCTGGA TATTATCAAC TCACTGGTAA CAACAGTATT COCAGAACCT ACTOMOGCAG COMOCTGAGA AGAGTTGAGG GAAAGTGCTG CTGCTGGGTC CCGAGCTGGG CGAGAAGTAG GGGAGGGCAC GAGCCGCCGC GGTGGCGGTT GCTATCGCTT GTATTTIGACT TOTOTTCTCA GCATTCAGAG AGCAGCGGTG TAAGATTCTG CTGTTCTCCC CGGATTIATC CATTITIACT GATGGTCGTG GTATTGATGG CAATTITIGTC CTTCCAAGTC AGGGTGTTT GCACTIGIGA CAGCAGTAIG CIGICTIGCC GACGGGGCCC TTATTTACCG TOCAGACOCO ATOGATAACO TOCAGOOGAA AATAAAACAT COCCOCTTOT GOTTCAGTOT GGTACCCAAA T TACCCTAAAA CCTTGGATTA AACAGAATOT GCATTGTACA TCTTTAAACA AAATGTATAT CTCGTGAACT ACGAACGGAA ATCTGGCAAA CAAGGCTCAT CTCCACCACC TCCACAGTCA GCITCIGGIG TIGHICCITY ACTAGGIGIT ACTGCGGAAA TGCAAAACTT AGTCCATCGG AGTITIGIAA TITTATATTA CITITIAGIT IGATACIAAG TAITAAACAT ATITICIGIAI (2) INFORMATION FOR SEQ ID NO: 98: TAATTTATTA AATCTAGTTG TCACTTTAAA AAAAAAAAA AAAAAACTCG AGGGGGGCCC TOGATICTICT GACATTACTO CIGICIGAGA TITOTATATO TOTABATACA AGTICCTIGA TGGACTICIC ICTITIGGAGA TITTITCCCAG IGATCICICA GCGITGITIT TAAGITAAAT COCCAGTTTA AGCOCCTTTA TGAACATATT AAAAATGACA AGTACCTTGT GOGTCAACGA GTAATTGAAC AGGITTACGC AAAIGGCAIC CGGAACAITIG ACCITCACIA TAITIGITCGI CTGCATGCCA ANATCATTGC AGCTATANCA TTGATGGGTC CTCAGTGGTG GTTGANAACT ICCCAAGAAT AAAGTAGTIG TCTCAACAAC TIGACCITCC CCTITACAIG ICCTITIITIG AMACTOSCAS CICCOGIGAT CICIGIOCIG TIGCITTICCC IGIGIGIACC ITAIGICATA (1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: <u>0</u> € (A) LENGTH: 504 base pairs TOPOLOGY: linear TYPE: nucleic acid STRANDEDNESS: double 98:

> 1620 1631

1560 1500 1440

1320

1200 1140 1080 1020

960 900

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480

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360 300 240 180 120

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.) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1416 base pairs
(B) TYPE: unclaic acid
(C) STRANDERNESS: double
(D) TOPOLOGY: linear Ξ

(2) INFORMATION FOR SEQ ID NO: 99:

9

SEQUENCE DESCRIPTION: SEQ ID NO: 99: (X

S 120 8 240 8 360 52 480 549 8 99 720 780 840 8 980 1020 080 1140 1200 1260 1320 1380 1416 GACAGGCTGC CCTGCCGGTT CCACAGGGTA CAGTTAGGAC TTGAGTCTTT CTTTTTCTGT TTTGAGTTGG TGAGTGAGTG ATAGGGTAAC ATGGGCCTTC AGGATGACCC CTTGGAACTG TGCCGAGTTC CITABATCIC AGCIGGGAIC CIGGACCIGG GAGOCCCCIG IGAGGGCCAG CICIGGAAAA CACTOGRAGE TETEGGGGTG GGAGGGAGAG GGGCTCCGGC TETETETGAA ATGAACACTG ACCTOGGAGT TGATGCCGGA GCTGTGGAAG AACTCTGCTC GAGGGCAGGG TGCCCTGGAA CICTICAGCA GITCAAGIAC ITGITCICAA AACAITITICI AAITGAITGG IAGGITITICA TAAGCATTOT ITCITITAAGG CATGGAAAGG GAAGAATOCT CAAGCAAGTC AIGITTIGITT TCAGTGGGAT GGCCCCCCT TCTCACTGCT GGGGCTTCC CCTTCATGTG GCACCTTTGT GCAGGGGCCA CCAGGCAGAC TCTTCCCACC TTCTCCCACT GAAGCACCAA GGGGCTTGGA ACCGINATIT GGCTANICAG AGGCATITIT TITICICCIAG INICITICAC ACTIGICCAA COGICITATI ITITIAAAG ITCIGIIGCI IGIATIAACA CGAAACIAGA GAGAAAIAGI ITCTGAAGCC AGTITATIGT GAAGAICCCC AAGGGAAGT ICGGIAGAGA AAAAIAGIAA GCTGGTTTNG AAACTGACGA GGCCAAACAG CCAGGACGCA TTGGAGAGGA ATTTGCCAAA GATCTACCCT GAGATAACGC CTGTCCAGTG TCTTCACCAC GTGAATAACC AGGGCTCCAA TTTAAGAATC CATGTGACTY TAGAATGGAA CTGCCGGCCC TGGCAACTGT CACGTGTGCT AGAAGGTTCG CTITITAGCAG ATCTGTCCCT GTGGGTGGTG TCTAAGAAGT CGGACACCTT GGTTTTTGTG THAGATTGAG CTGGGCAGCT GCAATCAGCT TCTTTATATG CAAATTAGGC ACGACCCATC IGTOGITICCT GÓTTGGTGGC TAATGAAGTG AGGGAGGGA GGGATGTCAC CCCAAAAGTA GCCCTCCCA TIGGCTTTGG CCAGGCCAGA CACITCACAT CGTTTACATG GTTCTGTGTA TTATGTGTAT AAAGCGAAGC TGTTTCTGTG AAACTGTATA TTTTGTAAAT ATGCCTCTGG AATGCATGTG ATACTCATCT CCATTITIGIT TCCTTGATTG CATTITITIGIT GGCACGAGGG AGGGAGCCCT CTCCCTTGGG TGACTCTTGT GTGCCCTTTA AGTOTITITIC TOCTITIGNAA AAAAAAATTC CACAAGCTTT TAAAAGGTGCA AAATATIIG CTACITGAAA AAAAAAAAA AAAAAA ATTITIABAGT 5 8 25 30 35 6 8 5 S

INFORMATION FOR SEQ ID NO: 100: 8 S

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2847 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double 3

TOPOLOGY: linear

2

SEQ ID NO: 100: SEQUENCE DESCRIPTION: ž

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140 260 980 200 GGCTAGGACA ATTITGGTGC TTTACCTATC TCTGCAAAGA CTGGAGAATT TGGCATACCA ICTCGAAAAA ATITAAGTAT CAGAAGATTA AAAAGAITTT AGGATTIGGA AGCTIGIAIT CAGTGGCCAC GICTITICCCC AATAATCAIT GITTGAICTC CAAATAGTAG CCITATAITA GCAATRGACA GATCATTICGT TOTCCATATIC TICATCATATIC TTACTACTITT GGAATCAGTA TITTGGGCAAA ITATICAATIOT AAGAACTAGG ATGCTTCCTG CAGTGGCACT ACCTTCCCCT AGACCTGGAG ITCAAGCAIT TATGCAGTGG ATATAAATGG AAATATAAAA ATATTTGCCA ACCTGTCTCA GRANCTIVATE APATETICIET GNATECTICAA GRANAGEACT TITICCTITTIA CITTAGAAAGE GITICAGAIT IGCITIATAG ACTOCTOCTG TCTTCAGIAC CIGATAAAAC ITTAACCAGG GAAGCATTAA ACACAGTGCA GCAGCTTTTG CCCAGGCTTC TAAGTTCCTG CCGGCAGCAT **GTCAGTAAAA** CTGAAACCCC CKGATACTCA CACTITICATE CTGACAGCCC AAGAAGGGAA ATAACTTGTA TTAAGGAACA ACTATGAGCC AGGCCCTGAG CTGTCTCTTA GATAATAAAA CAGATGGGGA GTGGAAGAGT CATTTGCTTC AAGTTATACA GCTAGGAAAT ACTICAAGCCA AATCTTGAAC GCAGCTCCCC CTAATTCTGT GGACAGGCAC TTTGTACCAC ACACCATGGT CCACCTAAAA ACAGAAGGAT AAAAAGACTT CAGGTTTTCC CACTGTGTGC TGACCATCCC AATTTATGAA TCTTCTTCAA AATGACATTT CACAGTTATA GTTAGGGCTC AGAMATOSCA TICAGGIAGE CITATITICIE CECTITAGEA GATOCITIAA GIACACATIG THANTIACAA CCACCAATCA TATCCAACAA AAGTACCCTA AAAGAAGGAC OCAGGGTTTG CTCCTGCCAG CAGAATCAGT TAGGAATTIC IGGCTAICTT ICAAATGITG AATTICIGGA IGCTGAGAGG IGATATCATT AAATCCAGGA CAGTCCCAAG AAGTGCTTGG AGTCTCGGCT CATGCTGCTT GOCCTTAAGC CCCAGCATGA TGAGGCTTCC GITTAGAGAGC TCAGAATTGG GTCTTGCCTG GGTGCAGGTG TAAAGAGAAG TCACCAAGGG AGGCAGGTAA TGAATGTTTC 2 2 25 39 35 45 25 **4** S

320 380

AAATAAGGTG TITCTTGGCC TICAAAGATA TAGAACTITIG CAGCAGTAGT AAAAGTGAAG

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CTGACTTGAG CCCACCCCA GGAGTTAGGA GAACATTTCC TTTTTTCATGC CATCTTCCAT

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TOPOLOGY: linear STRANDEDNESS: double

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 794 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

INFORMATION FOR SEQ ID NO: 102:

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Ξ SEQUENCE CHARACTERISTICS: 9 ۶ LENGTH: 1394 base pairs
TYPE: nucleic acid

INFORMATION FOR SEQ ID NO: 101:

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8 3 6 35 8 25 20 15 5 Ś GCATTGCTGC ATCCTGGGAA GGGGGTATAT GGTCTCACAA GTTGTTGTCA TIGTTTTTTT TOTOTOCAMA ACTOCAGAMO CTCACTOCCT ATAMGAGGAA ATAMGAGAGA AAGTOGAAGGA AAAAAAAAAA ACCC GCATGCTITC TTAATAAAAA AAAAAAAAAA ATGTTTANAG TTITIATCTTA AAAAAAAAAA GICCCIOCIC CICAIGITIC IGGITIOGIG AGICCITIGI GCCACCACCC ATAAIGCTIT GAGGGACAAA AGGAGTAATT ATTTGGTATA GATCCACCCA TCCCAACCTT TCTCTCCTCA TITICTIGTIG GICTAGCATI GCIGGACACA AAGIGTAGIC ATTATIGTIG TATICGGIGA CONGACTORS THIOGOSOMIC AGITTICCOMY COCCUTICATIG ARANGARARG RATACTROTT NTONOTORGA TROURGOTOS CTRUCACURA GCOURGITOTT GUTACOTOGO ACROGOURRA TICIGGCOIG CAGCAGGGIA GCIGAAGITIT GOGICIGGGA CIGGAGATIG ATCATACATO CAAATOCCCC TIOTICATCI OTGICIICIG CAAACTAGIC ICAIGAAGAA GGGTGTGTCC TATACAAAAC TTCCCATCAG TTCTCCTCAA TATTCCCCAT TTGTAAATGA AMAGAMATOG GOOTGCGAGT GOCTTGAATC TCCCATGATG TTGGAGGGCA CTTAGTGGGG GAACAATTIT ACTICIGICC TIATITICACI IGCIGAAAAG CIGIOGGACA AAAIGTAIOG TCACTTCTCT AATGCCTITG CCCAGTGTCC CTATTTTAGG CATCTTTTCC TTCCTTATTC CTTCCAGTCA tticaagtatg acataatatt teeexittogg gaaaggagaa titieteitag agggtogeaa AAAGGGGCAA GATAGGGCAG TIAACTAAAG AGCACTITAT TICITIGAAG CCITICIAAG CTITITITIGG AAGGGGGTTA TATATGAGAG TICATIGAAG AAGTCCAGIG AGGCIGAAGT ATAATTICCA AGAGGITIKG COTTCCGCIC TCCTGCTITT TICTTICATC CACCCCTITC AATAGACAAG GCCACTTICT TIGIGATTIC TGCTTTTCAT GCATATTATT TIATTIACCC GTTGCAAAGT ACTCAGTGTA TATTTTAATGT TGATTGTTGA ATTTTTAGTTA CGAGAGGGAA CCCCCAGAAA AAGGTATGGA GCTAACTCAT CTCTTTTACA AGGGGTGGCC ATGACTTACT GAGATTOOTG GAGGAGAGTA AATAATCTAG AGGCAAGAGT TCAGTGAGGG CCAAGGGGGA TITICTAAACC CITITICCTOT TCAGATCCAT ACAGGATTIG CAAGGGTAGG GCCATTAGGC 1380 1394 1320 1260 1200 1140 1080 1020 960 900 840 780 720 660 600 540 480 420 360 240 300 180 120 8

3 35 30 25 20 8 15 5 TABATTATTT TOTANGAGAG ATTTACTOCT ATCCCAGGAT GTTCGGACTT GGTGCCCCTG GGATACATTG GTGCAAAAAA AGCCACGGGS CCCATACTGG GCTTGATATG CCAGCAGIGG TIAIRAAGCC CCCTACCCTG TCCCATTCCA GAAACCATAA GACTCAGGCA TGCTGATCAC ATCAGATTIT TATGITITAAA AAAATCTCAT TATGGATTGA GTCCAGCCCA CATATAATGG CTGTGCAATA CATGCTTCTC AATAAGAAAA TTAACTGCAT GTTTACTGTG CICCAGITAG ACCTAAGOGC ACAAATOCAG AATTCATGAC CITGIAGITG TGGCAGGGIC TARGOCACTO AGTICARAGTO ACROCCCTGA AGGAMATTOC ACTICCAGCCC TCCTCCAGGA GTTGGGGCTT AGGTACTIGC TTACAGGAAG AGCAATICCC TAGCAAAGGT CATTAGCTCC ATTATICTICC GTACTACTIC AMMINITICT GICAGCCCTA ATTACANGIG TCACCATATA CATOTICCACA CACACACACA CAATATTTIGA GAGCTAAGGA AAACTICAAAG CAGCCCCTTC TCAAAGICCA GIGAAICIGG CICICITACI GATICCIGGI TITAGIGIGI GIGICGGGG TGAATCTITC AAGGGAATTA CACGIITGGG TTAATGITTC AGTATAICAT TITCATACTG GACAGCAGAT TAATACITAA TGAGGGTTAA ACCTGACCAG TCTTTCTACA GTGACAGGCC GTICTIGATI CIOGAGGCCI GCCIGGIAAG AIAAGAIAGI AIAATITIOGA ACIGAGAACA GCTCTAAGAG AAAAAGAAGG CCCATATGGG AGACTTCAGT CTCATTATTA TTOCCTTTAT TOTOTAATAA GATGGGAAAC TTGGATGCCC AGCCATTTTG GTGACCTGAG AGTCTAACTA AGTGTGTACC TATATATAAA GGACAAGTGT GATATGTGTG TATATGTATA TACATACATA NAMAMACTO GAGGGGGCC CGTACCC TOCATTTOGA AATCAATAAA CTATTACTOG AAATGCCAAA AAAAAAAAA AAAAAAAAA AGICACATIC CCICCITAGO AATCITCCCC TICCACCCIT TACAITAMAC AAGGGAACAC ACACTOCATO AATOGOGAGA ACCAATGAAT CCATTOTCCT CTOCCTATTT TCCTOTOCAC TACCAGAAAC AGCAGAACGA GGGCCAGAGC AGAAAAATGA AAATAAGTOG AGACACTTAT AGGAAATCTT CATATTTIAG TAAACTTAGC CGCCAGTGTA CTCTGTGAGG ATGTGGCAAT TCTCTCCCCA AGTAGAAAAT ATTCTCTTGC CATTCCTGAA ATTCCACATT ACTITICAGG 2160 2100 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2220 2040 1980 1920 1740 1500 2847 1800 1680 1620 1560

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:

101:

PCT/US98/04493

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1860

323

GGTGTTCTGC TCTCTACTCA ACTITATITIG AAAATGTCTG CAGCTTCACT CCTGTAGAAA

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(D) TOPOLOGY: linear		ACTIVITY OF THE PROPERTY OF A STREET OF A
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:		
GENRICAGOC GCACTAAAGG GACTTGAGCG AGCCAGTTGC CGGAITHATTC TAITTTCCCCT	09	S management introduction interchance whereasters instance were transfer of the second
CCCTCTCTCC CGCCCCGTAT CTCTTTTCAC CCTTCTCCCA CCCTCGCTCG CGTACCATGG	120	THE PROPERTY OF THE PROPERTY O
COGRECOTICE GCGOCACTE AGRECCATTE CATETECTES TESTECTIVES GAGGEGAGE	180	() ANACOTOR DESERVATION CONTRACTOR DESCRIPTION CONTRACTOR CONTRACT
STCCOCCCC GGCGGCGGC GGAGCCCAGG AGCTGGCCC GCCCTGGGGA CGAAGAACTG	240	
CAGCTCCTCC TOTOCOGTOC ACCATCTGAT TTTCTOGAGA GATGTGAAGA AGACTGGGTT	300	CHICHCERES ACTIONS ACTIONS ATTRIBUTED AND ACTIONS AND
TOTCTITIOGA CACOCTIGATC ATOCTOCTITT OCCIOGOAGE TITICAGTOTE ATCARTISTICS	360	לוסומסטיבע מוסומטיבע ווייניים מוסימטיבע אוייניים מוסימטיבע פווייניים מוסימטיבע פון
GITICITAMC TCATCCTOSC TCITCTCTCT STCACCATCA RCITCAGGAI CTACAAGTCC	420	OCCIONACIO GIOLOMACIO MOSTICIONE GIOCIONAMO CALIAMINO I CAUTICIO INCLUENZA I
GTCATCCAAG CTGTWCAGAA RTCAGAARAA GGCCATCCAW TCCAAAGCCT ACCTGGACGT	480	20 monthight managenag montharmas congagnam memorage cargadaes
AGACATTACT CTGTCCTCAG AAGCTTTCCA TAATTACATG AATOCTGCCA TGGTGCACAT	540	
CAACAGGGCC CTGAAACTCA TTATTCGTCT CTTTCTGGTA GAAGATCTGG TTGACTCCTT	009	יייייייייייייייייייייייייייייייייייייי
GAAGCTOSCT STETTCATOT GOCTGATGAC CTATOTTOST GCTGTTTTTA ACGGAATCAC	660 25	
CCTICTAATE CITGCEGAAC EGCECATTE CAGEGECCG AFFGETATG AGAAGTACAA	720	GAAACTUSS GALACATUTG GCCCTTTGCT TCTGAGGAC TCCCAGTGCC AGGCCTTTGA
GACCUAGATT GATCACTATG TTOGCATCGC CCGAGATCAG ACCAAGTCAA TTGTTGAAAA	780	
GATCCCAAGC AAAA	966	
	•	ATTICAMENT CICCITICAMO MAINIGAGGO TITITAGGATO TITIATATICO TICATOCCTO
	\$6	TIGITICCCA GOTTIGGAG GGAAAAAAG TCIGGAATIA IAGAIACAGC TIATIATIAA
(2) INFORMATION FOR SEQ ID NO: 103:	i	ATTIGITCIT GCATAAAAA AAAAAAAA AACNCANGGG GGGG
(i) SEQUENCE CHARACTERISTICS: (A) LENOTH: 1544 base pairs		
(B) TYPE: nucleic acid (C) STRANDERNESS: double	0,	(2) INFORMATION FOR SEQ ID NO: 104:
(D) TOPOLOGY: linear		OCTABLISHED SOMETHINGS (1)
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103;	45	3
TITIGETIGET ACTEMBACE ANABACTIOT TIGOSCATTI GETSTOTIGS CEATITICIGS	09	: © <u>@</u>
AGCAAGAGGG TETTETTECT CETTCCCCCA GCCAGCCAGC TGTCCTGGGG CCAGGCTTTC	120	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:
CTOGGTGGAA AGAMGTATAC CTTTCCCTGG GCCCCTAGGA TAGCAAMGTG AGCCATAGTG	180 50	ACCCACG
GOCCAGACTIC COCTOCATIOC TOGGOCOCIAG COCAGATOTIC CACTOGOCITE GATCACCTTIC	240	TIGGOCCCGA CGCCTCTGTT CTCGGAATCC GGGTGCTGCG GATTGAGGTC CCGGTTCCTA
TITICAGCCIT ACCUITCTCC TOTCAGGIAG GAATGAACIT OCCAGCCITIC AGGYTCGITIC	300 55	S AGGIGGGIGS CIGICCACCC GGGGGGGGG GAGIGAGGTA CCAGAITCAG CCCATTTTGGG
AGCTATGACC ATCTGTGGGG TCAGGGTACA CTCAGCTCTC CTCCCCAACT CCAGCAGCCT	360	
TTAAGAAGTG TCCCTTTCGC GCCCCCTCGA GGCAGAGCAC TGAGCTGGAC CCTGGGTAGA	420	

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 CTGCAAGATG GAGGAAGGC GGAACCTAGG AGGCCTGATT AAGATGGTCC ATCTACTGGT

CTCCCACAGG GAGGAGGAG CTGGCCTCAG GAGTGGGACA CCCAGACTTG GCAGGGCTT

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CT/US98/04493	

AATGCTTCTT CAGAAAAAAA AAAAAAAAA A TICCOCTACE AUGGOSTOTE CICICITIGE AATOTOGGST GEGIECIGAG CAATGGGSTE GATECITAAC GECAGNIOCO AGAGAAGGAC CECAAGTACA GIOCIETECG CEAGAATTIC AMACCOTOGO AGAAGGAGCO AGGCCTOGGT GGGGAGGTAC CAGGCAGCCA ACAGGTTCCC TOGCCACTOT CAACGCCCOC TOGCTOGAAC CCCGCACCAC AGCTGCCATG TGGGCCCTOC GOSCIPAGET CACATICIOS GASSCCASCE ASCITIACEI SCIVITECTO ASCETIACEC TECACATETE CATOGGETOT GECTTCATCA ACCTETIGAT CTTGGETTCA CAGCATGET TOTETEGETG GEETTGEEET GGAAATAAGG AGEETETAGE ATGGGEEETG CATGETAATA TOOGRAGOOT TOOOTGROAT ACCITOGGRO TAGTGCAGAG CAAACTOTTO COOTTOTROI 871 840 780 720 660 600 480 420

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(2) INFORMATION FOR SEQ ID NO: 105:

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(i) SEQUENCE CHARACTERISTICS: 3 <u>6</u> 0 8 TOPOLOGY: linear STRANDEDNESS: double TYPE: nucleic acid LENGTH: 404 base pairs

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SEQUENCE DESCRIPTION: SEQ ID NO: 105:

35GCCACGAGTT ATACCATIGC ATTICATACTT TTGTTTTATT GCCTCATGAC TTTTTTTGAGT

X

45 8 AGATACCAGA ATOGOTTACA CATTINACCT GOCAAACATT GAAGAACTCT TAATGTTTTIC TTAGAACAAA ACAGIGCAAC CGTAGAGCCT TCTTCCCATG AAATTTTIGCA TCTGCTCCAA AATTINTAAT NIGAAAAATT TITTIGGGGT TITTIGGGGCC ATGG TCTAAAATTT ATTTTTTTAA AAAGAGAAAC TGCCCCATTA TITTGGTGGG GTTGGTTTTT AACTOCTTTG AGTTACTCAG AACTTCAACC TCCCAATGCA CTGAAGGCAT TCCTTGTCAA TITITAATAA GAATGACGCC CCACTITIGGG GACTAAAATT GIGCTATIGC CGAGAAGCAG 240 404 360 300 180 120 6

(2) INFORMATION FOR SEQ ID NO: 106:

50

3 SEQUENCE CHARACTERISTICS: S 8 0 8 LENGTH: 1542 base pairs

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TYPE: nucleic acid STRANDEDNESS: double

TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:

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CTIGICAGGI GCCIGGGGCA IGCAAAIGIG GGIGACCIIC GICICAGGCI IICCIGCIII 360 20 2 5 CAGCCCTGAG GCACTTAGGA TOGTTCGGGT GGCCTGGCTC TTCCTCTTCT CCAAGTTCAT GATOGAGGCT GTTGTGAACT TGTACCAAGA GGTGATGAAG CACGCAGATC CCCGGATCCA ACATOTOTTO CATCACTOTO TOCTTOCOTO GAGOTOGIOG TOGGGGIAA AGATIGOCCO TRAGETGATG GACACAGTGA TETTTATTET CEGAAAGAAA GACGGGCAGG TGACETTEET CCTGATGTOG GOCTGOCTGA GCACCTATAC CTGGCGCTGT GACCCTGTGG CITCATGATT GICTACAACT TCTCACTGGT GGCACTCTCC CTCTACATTG GOOCTACCCT CTGATGOOGT CCCCCTTGCT AATGACCTCC ATTCTCCTGA CCTACGTGTA GOCTICTRICGA GAGGAGAGAG GAGAGCOCTO GAGAGGAGAG OCTOGAGAGT CCTTAGCICAG GTCAGACAGG TOGAGCCGCC GOOGCAGGAG TCTCAAAGAG CCAGGCTCCA GGAGAGGAAG

TCACTTGGGC CICGCATCAT GCCTAATCGG AAGCCCTTCC AGCTCCGTGG

ACTATTCCA TCTATGAGTT

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GGGGGGCCCG TACCCAAATC GCCGGTATGA TCGTAAACAA TC

AACTTOTOTO TTAATTAAAA GTGACAGAGG AAACCANAAA AAAAAAAAAAA AAAAACTCGA

1500 1440 1380 1320

1542

25

AATAATOTCA CTOTOGACCC AGTCTCACTC CTCCACCCCA CACACTGAAG CAGTAGCTTC

TGGGCCAAAG GTCAGGGTGG GCGGGGGCCT GGGAATACAG CCTGTGGAGG CTGCTTACTC

6

CCACTITCCCC COGGCAGCIC CAGOGATGTO OCCTCATTGC TGTCTGCCAC TCCAGAGCTG

GGGGCTAAAA GGGCTGTACA GTTATTTCCC CCTCCCTGCC TTAAAACTTG GGAGAGGAGC

1260 1200

ACTICAGGGOT GGCCCCACAA AGGGTCTCGT GGCCTTTTTTC CTCACACAGA AGAGGTCAGC

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TOOGTCAGTG COCTOTOCAC CTACACCTGT GACCAAGGCT TATGTGGTCA GGACTGAGCA

AGAAGCATOG CCTAGATAGG COCCCACCTA AGTOCCTCAG GACTGCACCT TAGGGCAGTG

GOGGACTOGO COTOCOCTOS COACAGOTOS TOTACAGGGA COACGGCTTT GOTTOCTOAC

1140

1080 1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 8

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CACCATCTTC TTCATGCTGT TCTCCAACTT CTGGTATCAC TCTTATACCA AGGGCAAGCG

CITTATISTICC AGCISTAACT ACCAGTACCC AGTCATTATT CACCICATCT GGATGTATGG

GCTGCCCCGT GCACTTCAGC AAAATGGAGC TCCAGGTATT GCCAAGGTCA AGGCCAACTG

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CATGACAGCC ATTCAGCTGA TOCAGTTTGT CCTGGTCTCA CTGCACATCT CCCAGTACTA

GTACTACOGA TTATCTOCCT TTOOCCCTOT GGCACAACCC TACCTTTOGT GGAAAAAGCA GGGAGGAATG GGCTCTTTCC ATGCCATGAT AAACTCTTCC GTGCATGTCA TAATGTACCT

INFORMATION FOR SEQ ID NO: 107: Ξ

8

SEQUENCE CHARACTERISTICS: (A) LENGTH: 2327 base pairs (B) TYPE: nucleic acid

328

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	329		. 330
	(C) STRANDEDARSS: double (D) TOPOLOGY: linear		TITIACANTE GACAANGAA ANTETHACAG CENGAAGGCA GAGGTGCC CAGANGTAA
v	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:	•	AGAGACCTIC AGTATCAGCC CTAACTCTTC TCTCCCAGGA AGGACTTGCT GGGCTCTGT
n	GOTACCTCAN TOCAGTGAAA TAGTCTTACT GGAAACAAAG CCCTTTATCA AGAATAATTA	09	OCCHOCTOTO CAGOCCAGGC CTOTOTOTO ATCOTTTOTO ACGISTOCIA ATGGGAAAG
	ACTICICCY TITICITITIG GAGAGGIGCT TIGITICIGA ICGGACCAIT TCACTGCAGC	120	AGGGCTTTTT ACATCTCCTA AGGACCTGA TGCCAACACA AGTAGGATTG ACTTAAACT
9	AAGCAACACA GTATTCTRAG CAGAAGATCG GGACTTGAGG CCATGTTGGG GAGGGCCAGT	180	TTAAGGGCAG CATATTOCTG TACACATTTA CAGAATGGTT GCTGAGTGTC TGTGTCTGA
	RACATTATCT GEACTCTGGA GTGTGAGGAA TATGGACTCC ACTCTTCACT AINITCACAR	240	ITITICATOC TOGICATGAC CTGAAGGAAA TITIATTAGAC GTATAATGTA TOTICIOGIG
ž	CANTICACAC TICAGCAACA ATAGCAGITIT TAGCCCIGAT GAGGAAAGGA GAACTAAAGT.	300	TITIAACTIG ATCATGAICA GCTCTGAGGT GCAACTICTT CACATACTGT ACATACCTG
3	ACANGATOTT GTACCTCAGG COTTOTTAGA TCAGTATTTA TCTATGACTG ACCCTTCTCG	360	GACCACTCTT GGGAGTGCTG CAGTCTTTAA TCATGCTGTT TAAACTGTTG TGGCACAAGT
	TOCACAGAGG GTTGACACTG AAATTGCTAA GCACTGTGCA TATAGCCTCC CTGGTGTGGC	420	TCTCTTGTCC AAATAAAATT TATTAATAAG ATCTATAGAG AGAGATATAT ACACTTTTGA
20	CTTCACACTC GGAAGAGA ATTCGCACTG CCTCAAGAGA ACGIANGAGA CTYNGGCCTC	480	TICITITICIA GAIGICTACC AATAANGGA ATTIGNGACC TOTAAAAAA AAAWAAAAA
	AGACATOCAG TOGAAAGTTC GACGGAACTC TAGCATTCTC CATCCAGGG CTTGCAGTTA	540	ACTICAGGGG GOCCUGGTAC CCAAATOGCC GATATGATCT AANCATC
ž	TICTTGGAGA TCAATTGACA GCTGCAGATC TGGTTCCAAT TTTTAATGGA TTTTTAAAAG	909	
3	ACCITCBATGA AGTEAGBATA GOTGITECTA AACACTICEA TGATTITICTO AAGCITECTIC	C7 099	
	ATATTGACAA AAGAAGAGA TATCTTTATC AACTTCAGGA GTTTTTGGTG ACAGATAATA	720	(2) INFORMATION FOR SEQ ID NO: 108:
93	GTAGAAATTG GCGGTTTCGA GCTGAACTGG CTGAACAGCT GATTTTACTT CTAGAGTTAT	30	"
	ATACTCCCAG AGATOTTTAT GACTATTTAC GTCCCATTGC TCTGAATCTG TGTGCAGACA	840	(B) TYPE: nucleac acid (C) STRANDENESS: double
35	AAGITICITC TOTTCCTIGG ATTTCCTACA AGITGGTCAG CGAGATGGTG AAGAAGCTGC	35	(U) 10FULUES: LINEAT (V) CENTENATO RECEDENTAL, CENTENA, 100.
3	ACCOCICAAC ACCACCAACG TTCGGAGTGG ACCTCATCAA TGAGCTTGTG GAGAACTTTG	096	: PAT : TO : TO : TO : TO : TO : TO : TO :
	GCAGATOTCC CAAGTGGTCT GGTCGGCAAG CCTTTGTCTT TGTCTGCCAG ACTGTCAITG 10	1020	SOCIOLOGIA GEOGRAPHICA GEORGIAGOSTE CARECTOGRETT AGGGRANATE
40	AGGATGACTG CCTTCCCATG GACCAGTTTG CTGTGCATCT CATGCCGCAT CTGCTAACCT 10	1080 40	TICCICATOS TCATCATOS CITOCICATO COGATOTOS TOCAGOTOCO TOTOCOCOCO
	TACCAAATGA CAGGGTTCCT AAGOTGCGAG TGCTGCTTGC AAAGACATTA AGACAAACTC 11		ALACAMANGA TIGGGBANGGT CCAACATGTAG TITCCICTIGGA GGTTCTCGAA GATGCTCTTC
Ý	TACTAGADA AGACTATTIC TTGGCCTCTG CCAGCTGCCA CCAGGAGGCT GTGGAGCAGA	1200	STATEMENT TO THE CONTROL THE TENENT OF THE TOTAL OF THE T
}	CCATCATGGC TCTTCAGATG GACCGTGACA GCGATGTCAA GTATTTTGCA AGCATCCACC 12	0921	CHANTEMEN ATTUCKTURE CELECOGRAM GRANGTRACA GRAGOCOGOG GTACHOCTET
	CTGCCAGTAC CAAAATCTCC GAAGATGCCA TGAGCACAGC GTCCTCAACC TACTAGAAGG	1320	FAIRCIANCE MUNICHILA CLUMMACHAA MANGICCIAC AAAAGAACCO TCCAATAGAA
20	CITIGNATICIC GGTOTCITIC CTGCTTCCAT GAGAGCGGAG GITCAGTGGG CATTCGCCAC 13	1380 50	SECTION TO TRANSPORTED TO SECTION TO TRACTICATE SECTION TO TRACTICATE SECTION TO TRACTICATE SECTION TO TRACTICATE SECTION TO TRACTICATE SECTION SECTIO
	GCATOTICACC TOGGATACCT TTCGGGGGAG GAGAGCTT CCTCTCCTGC GGACTTCAIT 14	1440	CHARLENG CONTINUE AND AND AND AND AND AND AND AND AND AND
Y	GCAGGTGCAA GTTGCCTACA CCCAATACCA GGGATTTCAA GAGTCAAGAG AAAGTACAGT 15	1500	CONTRACTOR CONTRACTOR INTERCOLUTE CONTRACTOR CATTLOCAGE
3	AAACACTATT ATCTTATCTT GACTTTAAKG KKWAWRAMW KCTCAGNSRA TTATAHTTSW 15	1560	SACTIMETICS ASSESSMENT ATTENTIONED CONTINUES OF SECURITIES
	CMMRARGSH WIMAAWSCTK SWCCTCYWCC KSRSTGRWKG MRCTCTAGA AYTRGYRGAK 16	1620	CYTCHARGA CARACTER ACTIVITIES OF THE OFFICE COORDINATES TO ACTIVITIES OF THE OFFICE OF THE OFFICE OF THE OFFICE OFFICE OF THE OFFICE OF THE OFFICE OFFICE OF THE OFFICE OF
8	CHYYYKSGCT KAMGGAAKKS GGCASGAGCC AGAGACCTOC ATTGCTITCT CCTGGTITTA 16	1680 60	CTICHMANN CLAMITICA TICICITON, TITCHALALT TOTTUMANLA GGILLAGAG

2100 2160 2380 300 420 480 ATC GACAAATGAA ATTCTTACAG CCTGAAGGCA GACGTGTGCC CAGATGTGAA ITIC AGTATICAGCE CTAACTETTE TETECEAGGA AGGAETTIGET GGGETETGTG STC CAGCCCAGCC CTGTGTGA ATCGTTTGTG ACGTGTGCAA ATGGGAAAGG ITT ACATETEETA AAGGACETGA TGECAACACA AGTAGGATTG ACTTAAACTE TAG CATATTGCTG TACACATITA CAGAATGGTT GCTGAGTGTC TGTGTCTGAT IGC TGGTCATGAC CTGAAGGAAA TTTATTAGAC GTATAATGTA TGTCTGGTGT TIG ATCATGATCA GCTCTGAGGT GCAACTTCTT CACATACTGT ACATACCTGT IT GGGAGTGCTG CAGICITIAA ICAIGCTGIT IAAACIGITG IGGCACAAGI TA GATGICTACC AATAAATGCA ATTTGTGACC TGTAAAAAA AAAWAAAAA ICC AAATAAAATT TATTAATAAG ATCTATAGAG AGAGATATAT ACACTTTTGA AG GCGCAACAGC CGITCTGTCA GCTCTGGGTC CAACCGGACT AGGGAANATC CE TCATCATOST CITICETCATO CIGATOTICAS TICCAGSTOCO TETECOCOCO CG TCATCATCTT CCTCTTCGTC TTCCTCATCC TCATCCA GTTCTCGAAG AC ATGACCATTA CCAAAGGCAA AGACTGCTAC AAAAGGAGCG TGCAATAGAA 36 TGGTCTTCAT TGGAAAGATA CCTGGCCGCA TGACTCGATC AGAGCTGAAA TI COSTITITIOS AGAGATICAS GAGIGCACCA TOCACTICOS TOTOCAAGGG 36 TGGCGAAGGT CCACCTGTAG ITCCTCTGGA CGTTCTCGAA GATGCTCTTC SA ATCCCCATCC CCCGGCGGA GRAAGTGACA GGAGGCGGCG GTACAGCTCT AG GCCCGGTAC CCAAATCGCC GATATGATCT AANCATC xi) SEQUENCE DESCRIPTION: SEQ ID NO: 108: (A) LENGTH: 1062 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDMESS: double
(D) TOPOLOGY: linear 1) SEQUENCE CHARACTERISTICS: RMATION FOR SEQ ID NO: 108:

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8 45 55 50 8 35 30 23 20 5 5 S 2 ATAAACMSCT TCACAATTC TCCATGACTC TTATATACTG CCTCATCTTG ATTTATAAGC AAAACCTGGA AAACCTACAA ACAGAATCAG AGGCCATOGA TACTGACAAC TGATTTTGTCT GTTTTTTTTC TCTGTCTTTT CCTACATCTT CCTAAGCTTT TTAGCAGGTA TATGTTGAAC ACTICTGTTT CATGGTTGAG OTTANAMIC TATACTOGAC AGITACAAGA AATTACCOGA GAAAAGCITG TGAGCTCACC TITATAAAAT GIIGIAGIGA AGCCCACAAT IGACCIIKGA CIAATAGGAG IIITIAAGIAI AATTWOGSAC AAABTBGHGG GGGDTCCAAA CHTWVTCGHG KAAMTTCTCT WAARMATYTK AAAATCAATT TIGIATAGIT TATTICAATC TAAATAAAAT GIGAATITIG TIWWATTAAA AATAAGIGIT GIGGITTATC TAGAAAAATA TGGAAAATAT TGCIGITATT TIGAANGGAA AAGCCIGCIG TIGTICCACA TCICGINGCI GITTACATIC CITTIGIGGAG AAACAAGGAT TICAGIGIAG ATTITIGICIT TCITGAACIT AAAGAAACAA AIGACAAAGI CAGAACCTTG AMAGICCCAT GTTTCCTTTT GCCTGATCTC TOTTGATGGC ACTCTGGAAT CIGITIAAAI GOCCCCIGIT TGAACICICA AGCITIGAAG ACCIACCIGI TCITICCAGAA AATCACTAAA TATCITTIGCC TATAGGACTC CATTGAATAC ATTAGCCATT GATAATCTAC TAMARCACGA CCTGGATTTA ATOGTGGACA CATATATTAA ACTCTATACR AKTAMOTCAG GGAACACTTT GACAGACCAA AGGTCAAGTA ACTTGGCTTT GCTTAACATA AATTTTGATA TGATGAAGGT TGAGAATGAG CGGTATGAAA ATGGACGAAA GCGTCTTAAA GCATATTTGA CTGACATCAA GITTITITCCT AAIGIGIAIG CATIGCIGAA GGICCIGIGI ATICITCCIG GGAAACACAG GOGGAAAGAT ATAGAGCTTC COTCCACCAT CTATGAAGCC CTCCACCTGC GAAGTGACTT ACCCAATCCT GACACGCTGT CAGCTGAGCT TCATTGTTGG AGAATCAAAT TOGOTOTTAT GAATTCTTTG AMGAATATAT TTTGAAGAGG TOTOGGAGGA AGGAATACAT TOTTTCCAGT TTAAKTCATT TTAGACATAG CATTTATTAT CACTGTGGAT CTCTACTTGT CHMMTYYYMK RKTYGAYMYW YOCGWMCGAG AAAAAGCCGT AAGGTGTATG TAGACCACTT MICTYYCTAC AKAYRAYTCM SWAWITIGTOG AAARYWSSTA MIGMSWOCWKK TAMMIRITWCG CCTCAGTCAT GGGACAACTC AAATTCAATA CGTCGGAGGA ACACCATGCT GACATGTATA INFORMATION FOR SEQ ID NO: 110: (i) SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double (A) LENGTH: 1751 base pairs TITIGGTGAAG 2539 2520 2460 2400 2340 2280 2160 2100 1200 2220 2040 1980 1920 1860 1800 1740 1440 1380 1320 1680 1620 1560 1500 1260

AACTIAAAGA TATATICICA GAACAGCACC TCAAAGCICT TAAAIGCITA TCICIGGTAC 1080 1020

CCTCTGAGAG TTACTATAAA GAAACCCTAA GTGTCCCCAAC AGTGGAGCAC ATTATTCAGG AAATGAAACT CCCTGGGAAA TTCCGCAGAG CTCACCAGGG TAACTTGGAA TCTCAGCTAA TACAATKAAT YWTRRYTTSM KRMAGMYAAT CCGAAAYTGT GGMAAMYAAA CTTGATATTC 960 900

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ACAAGAGCCT TIGGGAAAAA CYYCMAGGG CAAACCICIG AIGICTICIT IGCKMMSRI ARMTTITGAY ATRIMARYACT RIMTKSAYTY AAYGRIGTGA CWSGAWAATA TTRAASTYTA 840 780

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TAGTGACACA AATATTAGAT GGAATAACTA TATAGCTGGC CGAGCATTTG TACTCTGAGT GCAGIGICAG ATTITIGATIT CATIGITACT ATTGITGITC TIAAAAATGI CCTATCTITI

> 720 660 600 540 480

TOCTITITICAA AITITIAGIOG AACICCIGCA AOCACTIVIT TIAITOTITAG AITOGTATAAA TAAAGAAAGG GGTAAAGAAC TGAACGAAAT CTGCCATTCT CAGTGGACAG GCAGGCATGA

2

TOGATOACOA CAACTOCTTT TAGAACTIGA CAACGTAATT ICIGITICTTT TICAGAACAG CCYGKRONIGS RRGIAWYISK IGCAYKAGGG AACAATIGAG GAAGITIGIT CTITITICCA

8

YWAWCKGAAC AMANKCYGSW CYTCCWSYGC SKTRRMKRYC GYKSTATRRC WARWKSAKYM 420 360

35

TOGATITICT TCCAAAATGA AAOTIGTIOC TICTAGACIT TYAAGMGGA TWKCCCCMAK AACTGAGAAG TGGGGATTAA ATATGGAGTA TTOTCOTGGC CAGGCTTACA TTGWCTCTAG

AGGCTICCTG CCTTATGAAG CCGATGCAGA AATTITGGCT GTGAAATTTC ACACTATGAT 90 240 180 120

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GCACCTACCT GTGTTGGTGA GGTTTGTTGA TGAATCTCAT AACCTAAGAG AGGAATTTAT

GAGAGACTCA CACTICITIT COATIAICAC TGACGATGIA OTGGACATAG CAGGGGAAGA

SEQUENCE DESCRIPTION: SEQ ID NO: 109: (A) LENGTH: 2539 base pairs (B) TYPE: nucleic acid TOPOLOGY: linear STRANDEDNESS: double

25

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2 INFORMATION FOR SEQ ID NO: 109 Ξ SEQUENCE CHARACTERISTICS:

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5 AACTAGAGAG NAAAAAAAA AAAAAAAAAA ATTTAAAAAA CT

S TATTITGITIT TITTATAACAG GTATIGAAAC AAGTIAACTI GCATICCIAT GTAAGATAGG AGGGGCTGAG GGGATCCCCA GTGTTTGGAA CATAAGTCAC TATGCAGACT AATAAACATC 1062

TATAAAGAAA TOGAAAAAAG TOAAATAAAA AATATOTTOA ATCAGATTTT TTAAAAGGGG AACCTCCTCC ACCCCCTTCC CCTACTCTAG GGGAGAGAGC TGCTAGTGAG ATGACTGTTT 1020 960 900 840

AACCICAGGA GGIAACCITG GGCCCTICCC IGCIAICCIT ITICICCTIT GGAGGIGCCC

780

33

•	WO 98/39448	PCT/US98/04493	WO 98/39448 P	
	333		334	
	(D) TOPOLOGY: linear		TOCAGTOGGC, ACTOCAGOOT GGGGGAGAGA GCGAGACTOC GTOTCAAAAA AAAAAAAAA	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:		AAAAAAAA A	
S	AGCATGAAGC CGATGGCCOT GOTGGCCAGT ACGGTCCTGG GCCTGGTGCA AAACATGGGT	N 60	5	
•	GCOTTIGGCO GANICCIGGT GGTGGTCTAC TACGTAITTG CCATCATTGG GATCAACTTG	G 120	(2) INPORMATION FOR SEC IT NO. 111.	
2	TTTAGAGGCG TCATTOTGGC TCTTCCTGGA AACAGCAGCC TGGCCCCTGC CAATGGCTCG		10 (1) SEQUENCE CHABACTERITY.	
2	GOOCCTOTO GAACTTOGA GCAACTTOGAG TACTOGGOCA ACAACTTOGA TGACTTTOGG	240		
	GCTGCCTGG TCACTCTGTG GAACTTGATG GTGGTGAACA ACTGGCAGGT GTTTCTGGAT	300	6 0 a	
15	GCATATCOGC GCTACTCAGG CCCGTGGTCC AAGATCTATT TTGTATTGTG GTGGCTGGTG	360		
	TOSTETISTICA TCTOSSTICAA CCTISTITICTIS GOCCTGARTIC TGGAGAACTT CCTTCACAAG	6 420	AARGITGIGG IGGIAGCAIT IGGGIIAAIT CIRAITAIAG AGICICIAAREE	
70	TOGGACCCCC GCACCCACCT GCTGGGACCC CAGAGGCCAC CTACCAGATG	480	20 CCATABACTA ATCCCABACA ACATTGTCTT TTTRATGTTG TAGTGBACAG CAGAGARTTT	
	ACTOTOGRAC TECTETICAG GGATATTETIS GAGGAGCECO GGGAGGATGA GETEACAGAG	5 540	CAAAGGACCT TGCTAAATATC TGTAAGACGG CAGCTACAGC AGGCATCATT GGCTACATT	
	AGGCTGAGGC AGCACCCGCA CCTGTGGCTG TGCAGGTGAC GTCCGGGGCTG CCATCCCAGC	009		
25	AGGGCGGCA GEAGAGAGA GCTGGCCTAA CACAGGTGCC CATCATGGAA GAGGCGGCCA	7 . 660 25		
	TOCTOTOCC ACCAGGAG GNAGACCT TTCCTCTGAC GACCACTNA GCTGGGGACA	720	TCATTOCITY TOSCINGOGE TESSESTITICS CARACTECIARE STREETS AND AND ADDRESS OF THE STREET STREETS OF THE STREETS OF	
9	GGAACCAAGT CCTTTGCGTG TGGCCCAACA ACCATCTACA GAACAGCTGC TGGTGCTTCA	30		
3	GOGAGGCOCC GTCCCTCCC CTTTCTTTA TAGCTCCTTC AGTGABATT CCCTCGTCGA	840		
	CTCCACAGGG ACCTITICAGA CAAAAATOCA AGAAGCAGGG GCCTCCCCTG TCCCCTGCAG	006	שנייניישונו שנייניישור הוויישור הוויישור הווישורים ביותר הווישורים הווישורים ביותר הווישורים הווישורים ביותר הווישורים הווישורים ביותר הווישור	
35	CITICOGIDOR OCCITIVOCTO COOGLACCO TIVOGGALCCA CAGGCCTGAC CAGGCCTGC	35		
	ACAGGITIAAC COTGAGTCTG TCTCATCTAT TCACAGCTGG GAATGATACT AATACCTCGG	1020	TABABUTICES SESTIMENTICES COCHESTINE ASCRESSIA COCHESTINE CONTRACTOR COCCUTANCE	-
Ç	ATTITIAGECE AGENECACAG GOTACOTICE AGITITICE TETTICEATA GENGTAAGGE	1080		
₽	CCTITICIOSO ANIGOTICIC ATTENCETTA ANCIAITATE GGOTCAGITE TECTGCATGE	1140		
	COCCAGOCTIC CCATCACTIGG CACCCACTICG CCACAGAGAT GOCCTGCTCA TOCGACTIGGG	1200	THE BELLING AND CLITCH VIANTAGATA AND AND AND AND AND AND AND AND AND AN	
45.	OCTITICACTE CCACACTOTO TACERCIETY GIGTGGACSE CCTOCTISCEA AAACTITCAG	1260 45	CANCITATION CONTRACTOR CONTRACTOR CONTRACTOR ACADAGAGG	
	CAARCAGCTT TOCAAATGGA AGTTGTCACT GTCAGGCCTT TACAATCAGC AACAGCAAAA	1320	CHARLES CHICAGO CALIGORY CALIGORY CALIGORY CALIGORY	
S	TCTACATGCT GCTGAGGGTC CTGCCTCATT AAGATGCAAT AAATATGTAA GTACATAAAA	081		
3	ACAGCAATAG AAGAAAGGTA ATGCTTTATT CTCAAATATG ATGTCTACAT AGAAAAGCG	1440		-
	AAATTATTAA GAATAGTAAG AATTCACCCA GCACTTTGGG AGGCCGAGGC GGGTGGATCA	1500	GAATTITACA GITAAAAAA AAAAAAAA AAAAAAA	
55	TENGCTCAGG AGATCGAGAC CATCCTGGCT AACAGGGTGA AACCCCGTCT CTACTAAAAA	1560 55		•
	TACANANAT TGGCCGGGCG CAGTGGCGGG CGCCTGTGGT CCCAGCTACT GGGGAGGTG	1620		
9	АОЭСАБСАБА АТОВОСТСВА ССООВДАЛЕС БОЛОСТТОСА БТОЛОССЬС	09 0891	(2) INFORWATION FOR SEQ ID NO: 112:	

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SEQUENCE CHARACTERISTICS

(A) LENGTH: 1313 base pairs

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(C) STRANDEDNESS: double (B) TYPE: nucleic acid

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 112

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8 25 8 35 30 25 20 15 5 СТТТТИННА АНАНАНАНА АНАНАНАН ЛАНАНАНА АНАНИНАНА АДА AATGCTTTTT TTGTTATCAG AGATTGTGTA CTATTTTTAT TTTTAATAAA TGTATCTTCC TITITITIAA TIAATATOTO TOCATTOTTA CAATOTATOT TOGGATOTOT TITICACCOTA AGAAGGAAGC CAAAATAGIT TITICCITIT GAAAGTITIT TAAAAAITAI TICAIGGGIC GACCACGIGA AAGGGAATOC IGGICTAGCI GGCGIGGIAI GITIHTAGGC GAATIICAGC ATTICAAAAAC AATGTGTTCA TCAAAGTAAT TGCTCACATT GTGCAGTACT ATGTTGTACA TITITICICAA ATGGAACCAT GGATTTATOT CIGGATCATC CATACAGAAC CAACAATTIT GTATTTGACA CTCATGCAAA ATAATGTGAA AACATCTAGA TTTAGTAGTT TATTCTGCGC CTITITOTTAA AACIGAAGAT TITIGGAAAAT GGITOTCACT GCICTICCAG CCIAIGAAIA GOTTATITAA GCOGAAGACT ACTITOCOATO CTCCAGGACA TGAAAAGACT GAAGATAATA TCAGTTAAGT AGTIGGTAAC CCTTTTCTAT TTTAGTAAAA CTTAATGCAT GTTTACTTTT TITIGATOIT TAGAGACACI AGITITIGGCC AACITAAGAT TITIACGITAA TITITIACATA TGACAAGCAT TAGTGACAAA GOCAGAAAAG ATTTATCAGC CATGCTAAAA GAGTGAAGAA TOGENATITE GGENCIGICE TOCCITCAGCA GEGCATATEC TETEGCAAAG TECTETGGET AACTITICACA GICCAGIATO CAACAGGAAC IGIGIGIGIG TIAAGACCGA AGTIACAATA TAAAACAAGC TTACACGAGT GCTCCAATGG TAGACAATGA ATTACTTCGA TIGAGTCTTC TACTACCAAA AGAAGAAAGA GCAAGICTTC TIAGIAATCT TOGCCCATOT TOTAAGGCG TOTOCTICAG ACOGGATITCT GCAATICGAA AGCAGCITTOT TAAAAATGAG AAGGGCACCA TITIGAAAGA TOCAGACAAT CICITIGAAC AIGAATIGGG GGCICICAAT AIGGCIGCAI TITIGAGACTA CCAACATAAC TACOTOTIGA AGGIOCTICA CAGAGAATAT ATIGCCITIA GCCAGAGGIT TICTTATATT TINAGTAAAT TIAAAGTGGC TATCAGAATA TITATTCTTG ATGTGAAATA ATTTTCACCA ATGTTGCTAA CITTAATAAA GTATAAAAIT TGTAGAATAT 1313 1260 1200 1080 1020 960 900 840 780 720 660 9 80 420 360 300 180 120

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9 SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 113:

(A) LENGTH: 1654 base pairs (B) TYPE: nucleic acid

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CARAAAAAA AAAAAATAAA NTTCGAGGG GGGC

1654 1620 1560 55

GATTTITATT CCCAGGATAT GOTSTICATT TTATGATATT ACGCAGGATG ATGTATTGAG

TAMAATCAGT TITGTAMATA TGIMAMINIG TCAUMAMIAA ACAMIGCITI GACITATITIC

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(C) STRANDEDNESS: double (D) TOPOLOGY: linear

20 5 5 OCCITICATOCT COGNACIONO TEMONOGONO MOGNAGANGA GAGANOGNICA GANGGAAGAG CAGIGGCAGI CCAIGGCIIG GITGAAGCIA GAAATITICC IGCCCCIGGI GACCIGGIAA CATAGAATCA CAAATACTGA CATTTCATTA GATGATTATT TTCCTAGAAT CCCCAAAGAG TICTOCTACT TICCCICCTA TIATAAGGAA ATTITACAGA TICTAAAAAT ACCITAATTI TCATGACAAC AACAGTCTTT CATTACAGAC TGAAGGGAAG CATGTCCTTA CTTAAAATAG COTCTACACT TTATTTTAAA AGCTATCCTT TTCTAGTAGT ATTTTATCAT GGCAATGGCA TICTITGATI TITATITITAC CAAGICACAA ATGICTITITI GAIGITITIGA GAATIGITCI ACAGGGACAG AATACTITICT TICCTITCCIT CAAGTACAAG AAGGCITICT CTACCATITIG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 113: 300 240 360 180 120

TICAAAATTA IGCAAGITAG TAATTACICA GGGTTAACTA AATTACITTA ATATGCTGTT GAATGCAGGA AAAATAATTT AATATCAACC TCAGTTGACA AGGTGCTCAG ATTATTCAAT CAAATTGATA ATATTCTATG TTCTAAAAGT TGGGCTATAC ATAAATTATT AAGAAATATG GAAYCTACTC TGTTCCTTGG CTAGAAAAAA TTATAAAACAG GACTTTGTAG TTTGGGAAGC AAAAGGTAGT GTGATAGTAT AAGTATCTAA GTGCAGATGA AAGTGTGTTA TATACATCCA TICTITATIC CICITICITC TGAAGATTAA TGAAGITGAA AATIGAGGIG GATAAATACA TICIGAAIGI TIAGGCAGIG CIAGIAATIT CCICGIAAIG ATICIGITAT TACTITICCIA TOGGGATECT COTTITICITA GOTTITICAG ACAACCOTAG ACCTAAACTG TGTCACAGAC ACAAATTATT GAATTTTATA AGCTTGTACA CAATATTTAA TTAGTGTGAA AGGAAACAAA TITIGIGATIT TITITITICIT CCGAAGAACT CCIGGITGIT AITGGATITI GTATITIAAT TCTCCCTCTG TCTTAGGAAA AGCCATCTTT AATATAGTTC TTCACCACTG TTGGGGTTGT CCCATTICCT TGAACCICCT GCICTAGCCT TGGCGGAGGG AGAGIGCIAT TIGCTTITGT AGTOTOAGGO CACAGITITOO TICACTAATO GIOCAGOTIG AGIGITOTGI TOTOITOCIG ATGGCAGCTT AGCCAGGTAG TCTTAGTGGT GGTGTTTAGG CATAAGATAT GCTGATCATC TOOTTAGATT TOOCTOTTOT AAAAGGGGCA AGAAAAGTAA CTCATCATCT CTAACACACC GCAGCOGANA INGACTIGACT TCACATOCTC AGCTITICTCA GCCITITIGIT TATTITIGITO 1500 1440 1380 1200 1080 1140 1020 960 90 840 780 660 600 720

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337		338
(2) TANDOBARATON FOR SEC TO NO: 114:	50	(A) LENGTH: 842 base pairs (B) TYPE: nuclaic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear
(4) INFORMATION FOR SELECTION SELECTION		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:
(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1171 base pairs		GGICTIGGGC GGAAGTGCAT GAGCTGCCGA TGTGGTGCTT AGTGATTGCG GTTTCGGTCG
(B) TYPE: nucleic acid (C) STRANDEDNESS: double	01	CTCTCCCGTG TITCCCGGGC TGGGTATTTG CCTCGCACCA TGGCGCCCAA GGGCAAAGTG
(D) TOPOLOSY: Linear		GOCACGAGAG GANGANGCA GATATTTGAA GAGAACAGAG AGACTCTGAA GTTCTACCTG
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:		COGNICATAC TGGGGGCCAA TGCCATTTAC TGCCTTGTGA CGTTGGTCTT CTTTTACTCA
GOCANACTIT CCCCCANGC ITCGANACTI GCANGOCGNA ACCITGANIC GITNANGIT	60 15	TOTOCOCTA THISSECT SETTING COUNTRANCE TRACARDERY TRACACOCAGO
GASTITICABNE GASACETTAS COCANAGANG COCANITICAC STICCOCAN COTTOGCCCT	120	ABCTEROTOR TOTAL TOTAL STATEMENT ASSESSMENT ACTIONS ASSESSMENT
CAACOGCITCO GCAGCCAGCC ATGTCCTGCA CCCAGGACAG CGGCCCTGGG CTACAAGGAC	180 20	CONTRACTO ACTIVIDATION CARACTEC ATTENDATES ACTIVIDATES
CTGGHCCTCA TCTTCCTGCG CCGACCTGCG CGGGGTAAGG GGHGTTTCA GACTGTGAAG	240	
GACCITCTIC TGGACTGCCT GITGGACTTC TTACCCGAGG GGGTGAACAA AGAGAAGATC	300	clearchacha iteathcaid deitheacha iteiteirie aidichean tirtiodeil
ACACCACTCA COCTCAAGGA AGCTTATGTO CAGAAAATGG TTAAAGTTGTG CAATGACTCT	360 25	CHARLECAN COCCARCOCT THACTICES TO SHAME INCIDENCE CHANTILLACE
GACCGATICAA STCTTATATIC CCTGTCAAAC AACAGTOSCA AAAATGTIGAA ACTGAAATTT	420	GCMGACATO GCACCCCANSC ACCAGANGCAC ANTENGAMAC GGCAGGGGGG ACMGGAGGGG
GIGGATICCC TCCGGAGGCA GITIGAATIC AGIGIAGATI CITITICAAAT CAAATIAGAC	480	CONCAGRIGA ACCOSTRIA GCCRITGACA TIGIOSCCAC AGGCCACTOS CCCTGGGGTGG
TCTCTTCTGC TCTTTTATGA ATGTTCAGAG AACCCAATGA CTGAGACATT TCACCCCACA	540	CICTICICAGE GICACAGC CCICATACCT GAACAATGA GAGICTAGT C CAGAGGCAA
ATAATCGGGG AGAGCGTCTA TGGCGATTTC CAGGAAGCCT TTGATCACCT TTGTAACAAG	009	ANCHAIRTIGG TAINCITHE CICHAINGG TOUTHWAIN ANTOCCTING
ATCATTGCCA CCAGGNACC AGAGGANATC GGAGGGGAG GCCTGCTTAN GTACTGCAAC	660 35	ARIGIGANA ANAMANNA ANAMANITO MOGOGOCO GOINCOANI ILUNCOIMM
CTCTTGGTGA GGGGCTTTAG GCCCCCTCT GATGAAATCA AGACCCTTCA AAGGTATATG	730	AT.
TOTTCCAGGT TITTCATGGA CTTCTCAGAC ATTGGAGAGC AGCAGAGAAA ACTGGAGTCC	780	
TATTIGGAGA ACCACTITICT GGGAATTIGGA AGACCGCAAG TATGAGTATC TCATGACCCT	840	(2) INFORMATION FOR SEQ ID NO: 116:
tcatogagtg gtaaatgaga gcacagtgtg cctgatggga catgaaagaa gacagacttt	006	(1) SEQUENCE CHARACTERISTICS:
AAACCTTATC ACCATGCTGG CTATCCGGGT GTTAGCTGAC CAAAATGTCA TTCCTAATGT	960 45	
GOCTAATOTC ACTTGCTATT ACCAGCCAGC CCCCTATGTA GCAGATGCCA ACTTTAGCAA	1020	(C) STRANDENESS: GOUDLE (D) TOPOLOGY: linear
TIACTACATT GCACAGGTTC AGCCAGIAIT CACGTGCCAG CAACAGACCT ACTCCACTTG	1080	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 116:
CCTACCCTOC ANTIAAGAN CATTIAAAA TOTCCTOTOG GGAGCCANT TCAGACAAGA	1140	GOCACOAGOC GOCOOCAGOG GTOOCOGOGG COCCCCCOG CGGGAGCCGT TCCCTTTCCC
CAGGAGAGA, AAAAAAAAA AAAAAAAAA A	1171	STCGGGGACC GCGGGGTCGG GGCCCAGGGG ACCCCGGGGCC ACGGAGAGCG GGAAGAGATAT
	55	GGATTGCCCG GCCCTCCCCC CCGGATGGAA GAAGGAGGAA GTGATCCGAA AATCTGGGCT
(1) semandaran and esp IP M2, 115.		AAGTGCTGGC AAGAGGGATG TCTACTAGTT CAGTCCAAGT GGTAAGAAGT TCAGAAGCAA
(2) INFORMATION FOR 30% IN THE STATE OF THE		CONTRACTOR CONTRACTOR

OCCICAGITIG GCAAGGIACC TGGGAAAIAC TGTTGAICTC AGCAGTTTTG ACTTCAGAAC

(i) SEQUENCE CHARACTERISTICS:

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CACACAAGCT CTGCGCCAAT CACAGGGGAA GTCTCCGCTG CTGTGGAAAAA GAACCTGCTG CTITHOTIGCA TCAGATGTAA CAGAACAAAT TATAAAAACC ATGGAACTAC CCAAAGGTCT TTTGGCTTAA CACATCTCAA CCCCTCTGCA AAGCTTTTAT TGTCACAGAT GAAGACTCAG TCAAGGAGTT GGTCCAGTAG CAATGATGAG ACCCTTTTAT CTGCTGTTGC CTTTATTGAA AGGGGACACC TGTACATTCT TCCATCGTCA CTGTAAAGAC AAATAAATGA TAGCACTTAC GTAAAACATT TGTTTCCCCC ACAGTTTTAA TAAGAACAGA TCAGGAATTC TCAAGCAGGA CCCTAAGATG AAGCTGAGCT TTTGATGCCA GGTGCAATCT ACTGGAAATG ATTATICAAA AAATCAIGIT TATTITGAGI CCIAGGACII AAAATTAGIC TITTGIAATA TAGAGCTTTT TAATAGCACT AACCAATGCC TTTTTAGATG TATTTTTGAT GTATATATCT ATTGAACAAA AATGTTTCCA CTGGCTTTTG CCTGTAAGAA AAAAAATGTA CCCGAGCACA GCCTAAGAAT ATGATCAGGT ATCTTOTOGO GAGCTOCTGA TACAGAAGAG ATGGATATTG AAATGGACAG TOGAGATGAA GAAACAGAAG AGCGAGTACA GCAAGTACGC AAGAAATTGG AAGAAGCACT GATGGCAGAC COMPRIMINGS METICSTMIKA YITHCATTICA ACTICIGATICE COGGGCCTTA GGTTTGACAT TAAATAAATT TCCCAGTTAA AGATTATTOT GACTTCACTG TATATAAACA TATTTTTATA TTATATTCCA CAGAAAAAAA AAAAAAAAAA MMSTYGARRR GSRGCMCRSW AYMMARWWCC 2 CTCACGAATT TOTOGIAATT TITTTTCCCT COCACCTICCC INFORMATION FOR SEQ ID NO: 117: Ê (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 117: AGGAAGATAG COCATATATT TOCAGTATGA ACTATTOCCT CTGGGACOTT GIGCTITICAC SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double g (A) LENGTH: 952 base pairs TOPOLOGY: linear AACTITICGAC CGACTITICCC CAAGAGAAAA TICCTIAGGAA CAGAATTITCT AAGGATTICT GGCTTAAATA TCACCTAGCC 1260 1200 1380 1140 1080 1020 1440 1320 1620 1560 1500 960 90 840 780 720 660 1640

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(2) INFORMATION FOR SEQ ID NO: 118:

E

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1256 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 118:

GACGTCATAG GTAAACAGGC TCTGTATCCG TGGCAGCGGC CGTGGCAGGC TGGCTGGGTA

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AICHTHOIMC CCCCAGCATC TAGCAGTOTT GOCATOTAGT AGGCACTCAA GAAATGIGTG GOOGCOGOG GACCGOLAGO TGAACTATOT GOTOTTCAGO GCGGGCACCG TGTTGCATTC COGCUTATION CTUALCICAGO AGANGETISCIC TOTICTACATIC ACCUTAGACTI GIAGOGGGCT

180

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25 20 5 30 5 S CATCTITIOG GTAAAGAGIT AAGIGICCAA AGGITGICAC AGITCAIGAG GICAGAGGA CTRAGGRUAG TETTITHAAAA CICCCAAAGA AAATCIGCIC TCCTTPCIGA ICTAAAAACT CTITITIOGIO AACTITAGIG GACTICIGIG AGATIGIAGI IGIACITIGI AICICIAAAI ATTITICCIOC CAGATACCIT TATICAAAATT ATTIOOCCICA TOAGAGCIGA AGTAAGICAG ATTITAATIT GCTATITTITI TCAATGITCT AGGTATCITT AAATTIGITA TIGIGGAATC ATTCCCTTTT TTGAAACAGG AAAAAAAATT ATTTTTTGTT CAGTAAAAAT GGTAGAGAAT THARTICTCC AGTAGACAAT GCIOGGTAAG GGAGGGGGTA GGOCTGGGTT ATTAAGATAG GUINOCCIOG CACCIGGACI CIGOCCATOC ACAGCIGACA GAITICCAACA GAAGIGIAIT TATCTCARTA ARACCCCACT GGRACTCCAA ARAAAAAAAA ARAAAAAAGA NN ATTYCAGAAA CATTGGGGGA AGGGAAAATT GGCTTYCTCT TAATTGGCAG ATGTTCCAGT AGOCTGCTGT ATTITACATT GOTTGTGGGG GAAGGGGAGC CTGGAGAAAA CAAAGTCACT COGAGICTOC TOAGITTATA AGGITCCAAA AATATOGIAA AATCITOGIT TITGITAAIT GOGSSGOOG GOCTCTOTT TIGTIGOGAT GIGITAIGIT GIAIGIACOC ATATAIGGAC TOCAATOTOO OTAGOOACAA GGGACCAGTT OCACTGAGAA GTGAACAGTG GGAACTCAAA 120 600 540 80 420 300 180 952 900 840 780 720 660 360

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CARTTITICAA ACAACCOGTA ACCCAAAGTC ACAAATCATC CTAGTAATAA AGTGAAATCA

CAATCAAAAT AAGGGTAAAC CAGACTTGAA ATACAACATT GCCAATTAGA CAAACAGCAT TOGRARGATG ATOCCTROTA ARTIACAGRA GRACABACAG RGACTGCGAR ACGRICCTCT

GACCCACAAC GAATGAATGA ACAGCCACGT CAGCTTTTCT GGGAGAAGAG GCTACAAGG

540

480 420

8

TOOTGACCIT CTAGATGAGE AACAAAAAA TGAAATAAGI TOITGGAAAT TAAGCCATIT

8

TGAATTTAGN AAACACTTTG GAAAACTCAT AACCTCATCA GAAACTGCCT TIAGCCACAC 60

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ATGACACAC TECTECTGAC TOCCACTOTE ACTECTTEAG AGCAGAACTE CTETAGGGAA

TIGANIGANC GAIGCCIOTG ACANGCANGC GGACTITATT CITICCIGAC CCITOCICCI

GAACTGAACA AGCTGAGGGT GITGGACCCA GAGGTTACCC AGCAGACCAT AGAGCTGAAG

CCTUGATIGGG AAACAGCCAT GGCCAAGGAC ATCCTUGGTIG AAGCAGGGCT ACACTITIGAT

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GAAGACTOCA AAGACTTTOT GGACAAAATT GGCCAGTTTC AGAAAATAGT TOGTOGTTTA	540	TITACAGCTA CTGAATTICT TATAAGGAAG GAGTOGITAG TAAACTOCAC TCTTTCTSTO
ATTGACCTTG TTGATCAACT TQCAAAAGAA GCAGAAAATG AAAAGATGAA GGCCATCGGT	009	arrangitas antagasena titragantes acoccoasto ecitosiscon Caasticoles
OCTOGRANCT TOCTCHANTE TATACCARA CAGAGARG CTCHACAGCA GCARCTTCAR	\$	THEATHER PARTICOL TARABELISC CICINITIAN PACACCION
OCCCTAATAG CAGAAAAGAA AATOCAOCTA GAAAGGTATC GGGTTGAATA TGAAGCTTTG	720	AGAGGGGCTT TATAAGCAGG CTGGGCAGG CCAGCTTATA AGTTAAAGGG CATCAGAGTG
TOTANACTAC AACAGAACA AAATGAATTT ATTGACCAAT TTATTTTTCA GAAATGAACT	780	
GABARITICO CITITRIROT ROGBAGOCIA PACABABAA AGCCICICAA BACCABABAA	840	
ACCICIOTAG CATTOCAGO GCTTGACCAA TCACCTATOT CACAAGAGGT GGCGTOTAAG	006	TITERARCEG ACTIRCATOR ARACOGRAPH CONCINERY CAROTHINIT ARACATIRAT
GANTGCAGCE CCETGAAGAE AGCACTACAA GTCTGGGGGA GCCAGTTTTA ACATCAGTGC	15	
ACAGCIGCIG CIGGIGGCC TOCAGIGIAC GITCICACCI CITARGCITA GITGGAACIA	1020	GAN
ACCAGITICS AAACTITCAT CCTTTTTTT GTAANTICAC AAAGCTTTGG AAGGAAGAC	1080	
ANIMAMITIT TETTITICAM TESCITICATE TACCITITIT CCTOTICCIC TICAMINIG	1140	
TITAACTOCT CATGAGAAA CCCTGGAITIC TCTATCCCCT AGTCCACAAA ACAAACCAGG	1200	(2) INFORMATION FOR SEQ ID NO: 120:
CAGTOSTCAG CAGCTACCTT INVITITIOSAT CACACACGTG AGTCAGACAG TACCAC	1256	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1782 base pairs (B) TYPE: nucleic acid (C) STRANERESS: double (C) PROMORESS: double
12) TATACHART OF SEC TO NO. 119:	30	
		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120:
(i) SEQUENCE CHARACTERISTICS: (A) LENCTH: 1143 base pairs		CAGGOCCOGG COCCCCACCC ACGTCTOCGT TOCTGCCCCG CCTGGGCCRG GCCCCAAAGG
(B) TYPE: nucleic acid (C) STRANDEDATESS: double	35	CAAGGACAAA GCAOCTOTCA GGGAACCTCC OCCGGAGTCG AATTTACGTG CAGCTGCCGG
(D) TOPOLOGY: linear		CAACCACAGG TTCCAAGATG GTTTGCGGGG GCTTCGCGTG TTCCAAGAAC TGCCTGTGCG
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 119:		CCCTCAACCT GCTTTACACC TIGGITAGTC TGCTGCTAAT TGGAATTGCT GCGTGGGGCA
восстатився всекваеттве тестветаеть маскваекае скажитает всякаект	60 40	
GIACCTICCA CATTGAGIAI TCAGAAAGA GTGATCTGAA CTCTGACCAT TCTTTAIGGA	120	TCTTGTTCCT GATTGCTTTA GTGGGTCTGA TTGGAGCTGT AAAACATCAT CAGGTGTTGC
TACATTANGT CANATATANG NOTCTONCTA CTTCACACAC TOCCTCGNOC ANACATGANC	180 45	
CITICARCITIS CCCACAGRAA AGTGAATCCCA AATACCCGTG TCATGAACAG CCGGGGTATG	240	
TOSCIGACAT ATOCATIGGG AGTIGGCTTG CITCATATIG TCTTACTICAG CATTCCCTTC	300	
TICAGIGITC CIGITGCITG GACTITIAACA AATATIAIAC AIRAICIIGGG GAIGIACGIA	360 50	
TITTICCATG CAGIGAAAGG AACACCTITC GAAACICCIG ACCAGGGIAA AGCAAGGCIC	420	CICCARICAL AGAGAARIAN GCNGBAGAGG THINGAGAIT TOTHOGNGGC ALTIGGCCTGT
CTAACTCATT GOGAACAACT GOACTATOGA GTACAGTTTA CATCTTCAGG GAAGTTTTTC	480 55	TUTICAGITI TACAGAGATC CTGGGTGTTT GGCTGACCTA CAGATACAGG AACCAGAAAG
ACANTITICIC CANTANTICI ATAITITICIG GCAAGITICI ATAGGAAGIA TGATCCAACI	540	ACCCCCCCC RATCCTAGE GCATTCCTTE GAFGAGAAA CAAGGAGAE TECCTTTCGE
CACTICATEC TAAACAGAG TICTETECTS AGTOTACTAA TICCCAAAAT GCCACAACTA	009	ATTATGATOT TOTTCACTIT CTOTTANGCT CCATTTGCCA GTTTANGGAA
CATGCTGTC GCATCTTTGG AATTAATAAG TATTGAAATG TTTTGAAACT GAAAAAAAT	09 099	

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25 20 5 30 5 3 8 છ S 50 TACTOGAAAA AGAGTOGRAA TTIATTAAAA TCAGAAAGTA TGAGATCCTG TTATGTIAAG CITAGEATIT TIACCIOCAG AAAAACITIG TAIGGIACEA CIGIGIIGGI TAIAIGGIEA GUICGOTOGO ACCIGAGAAT TIACIGIATI CATIGICAGA CACIGICCAC IGIGACCITI GGAAACACTA TCTOGAAAAG TACCTTATTG ATAGTOGAAT TATATATTTT TACTCTATOT GTAGCAAAAA GATATTTGAT TATCTTAAAA ATTGTTAAAT ACCGTTTTCA TGAAAGTTCT ANTOGRACGA CTTTTCAGTA ATCAGGAAGT ATATCTATAT GATCTTGATA TIGTTTTATA CAGAACTOCT TICATGAAAT CITICTAATOT ATAATAACAT TTACCTICAG CCTCCATCAG AAATAGTTAT GYCYTAGGAA ATTGIGGTTT AATTTTTGAC TTTTACAGGT AAGTGCAAAG TOTTAGTATA AAAATGATAA TIWACIKGIA GICTITIAIG AIWACACCAA IGIAITCIAG GGAAATCCAA ATTCCCAATT TTTTTTGGTC TTTTTAGGAA AGATGTGTTG TGGTAAAAAG ATCTGAACOT ACATCTCACT GOTATAATTA TATOTAGCAC TOTOCTOTOT AGATAGTTCC TICTCIACAT GITTITITCT TICCGTIGCT GAAAAATAIT IGAAACIIGI GGICICIGAA GARATTGARA TCOTATTOIG IGGCICTGIR TATICIGIIA ARRAKTIRAR GGRCRGRARG CAGTATIGTA ACAGCAACIT GIYAAACCIA AGCATATIIG AATAIGAICI CCCATAATII ATARTYTGAA GICTAAAAGA CIGCATTITT AAACAAGITA GIATIAAIGC GIIGGCCCAC TTOGATAGGA ATGGGGCTGA TGGGCTTCAT CGTTTATAAA ATCCGGGCTG GITGGCTGCA GATTIGIGGI GCGITCTGAG CCGICTGICC IGCGCCAAGA IGCTICAAAG (2) INFORMATION FOR SEQ ID NO: 121: CTTTCTTTGT GTATGCATGT TTGAATTAAA AGAAAGTAAT OG TATTATTAAA AACATATOGA TCCCCATGAA GCCCTACTAC ACCAAAGTTT ACCAGGAGAT AAGTAAGGCT TTGAAAGCTT CAGCGCCTGC TCCTGGTCAT CACAACCAGA Ê (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double (D) TOPOLOGY: linear (A) LENGTH: 610 base pairs TITACTIGGA CTGATAAAAG 1140 1080 1020 1380 1320 1260 1200 1740 1500 1440 1680 1620 1560 960 900 180 120 240 S

23 20 5 5 မ ઝ S CCAGGCACTC ACTIGIATIC TACTGCTCAA TAAACGTTTA TTAAACTTGA AAAAAAAAA CTGAGCCACC CATCATTOTO MAATAATTAC CTCAGTTOTA CAGGACTTGG TGATCAGGAT CAGGOTTCCA GGACATAGIC TGAGGCAAGA TGGAGGGTAT GAGGGGGCTT CACACTTCAC GENGETEGEE ATETTEGEEN ACATGETGGG COTOTOGETE TECTTOCTTG TEGTTETETA GOTACOCCTG CAGGTACCGG TCCCGAAATTC CGGGTCGCCC ACGCGTCAGG CCACGCGTCC 2 aaaaaaaaaa CCAGTATAGG GGCTIGCTTT TCTACTCCCT CCCCCCAATA TAAAAATATA GACTTTTTAA TTCATCCCTT CTACCCATCA CAACATACAA AGCAACTACA CCTGGATTTT TCCAAACAAC TCACTACOTO OCCOTCAACA ATCCCAAGAA OCAGGAATGA AAGTGGCGCT TTCTCCGCCC ACCCACOCOT COGSCCACOC OTCOGAGOCG AGCCOGACTG OTCAGGATGA TCACOGACOT TITTATITICC TCAGAGTCIT CCTTAATCCT ATGGAACAAG AAGCTGCCAC TGAATAGGGC INFORMATION FOR SEQ ID NO: 122: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 122: Ξ SEQUENCE CHARACTERISTICS: ਉ 0 (B) TYPE: nucleic acid (A) LENGTH: 526 base pairs TOPOLOGY: linear STRANDEDNESS: double 600 610 540 480 420 360 300 240 180 120 8

ຣ INFORMATION FOR SEQ ID NO: 123:

AAAAAAAAA AAAAANTTOG NOOGGGGCC GGTACCCATC CCCCTA

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ACCIGCAAATT COTOGTOCAG CTGTTCGCCG AGGAGTOGGG CCAGTACGTG GACTTGCCCA COGGETACGA GETGETEATE CAGAAGTTEE TEAGEETGTA CGGEGACCAG ATEGACATGE CAGICCCOGT GCTCTTCTGT TICTCAGTCT TCGCGCGACC CTCGTCGGTG CCACACGGGG ACCORDEGET COTTOCOCTO COCAACAGOG GOGOCOGOGO COCTOGGGCA TOTACCOOTC COGAAATTCC COGOTCGACC CACOTCOTCS GOGGAACATG GCGGCTKCGG E X SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 123: (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (A) LENGTH: 2081 base pairs

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180 120

300 240 S

S

GTACATGTGA AAGAAAACGT CAGTCTGCCT GTAAATTTCA GCAAGCCGTG TTAGAIGGGG AGCGTGGAAC GTCACTGTAC ACTIGTATAA GTACCGTTTA CTTCATGGCA TGAATAAATG

> 360 300

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420

480

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OCCITATIONS CATACICION GINGAINATI TOTCATOCAS COCATOCAAI CAGAAITCICA GATCHOTOAG ANGCACHGCT ACCIGGTACT GCTTICAGIG TOTICCCCCT CAGCCCTCCG

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ECCESSORIA GETRGACCEG COSTOROCTA GOGGCAGGC COGAGCCGCG GCGGCGGAGC GAAAGGAATT GCCCTTACTA CTTTGGGTTT GGTTTGCCCT TGGCTTTTCT CACAGCAATG CAGICCICAL ADALDAICAG TOGCIGCCIT INCICIANCO ICITICCITIT AFFICATUATIC TIGGIGGICT ICTIAAGCAA CAGACICTIC CACAAGACAG ICTACCTGCA GICGGCCTG AGCAGCICTA CITICIOCAGA GAAGITICCCI TCACCICATC COTCGCCTGC CAACTGAAG OCTACTICIAG GICACTICAGT TGCCTGCCAT CCAAAGGGA TGGCGGGAT TGGAAGAAGC TOTOGCAGET CITITICECTO TICACETECE GECTGCCAGG GAAGGCAGGA CECGCTICTGE CAAGGGCCCT CTGCGTATTC CCTTCTCTCT GAGGAATTGA AATTTTTGTC TCTGGTGCAC IGROGATICET TCATGATGAG AGAITITGGG ACACTICETET CTCCTGTGTG TAGITGATAG IGCITICAAT AICCITGOIT CATAAAGGA AITICAAAIGC ACCAGGGGIT GICIAACAIA ACCOCCAATG AAGCAAAGAC COCTGGCAAA GCRTATCTCT TCCAGTTGCG CCTCTTCTCC GIAAGGCAGA AIGIICCCIG ACACCAGIGI GIGGAITIITI AACAICACG IGAGICIGAA ITTIGGTGGTG AAGAGATGGC TGACAGTGTC AAAACCTTTC TCCAGACCT TGCCAGAGA ATCAAAGACT CCATCTGGGG TATTTGTACC ATCTCAAAGC TAGATGCTCG AATCCAGCAA AAGAGAGAGO AGCAGCGTCG AAGAAGGGCA AGTAGTGTCT TGGCACAGAG AAGAGGCCAG agtatagagc ggaagcaaga gagtgagcca cgtattgitta gtagaattitt ccagtgitgt GCTTGGAATG GTGGAGTGTT CTGGTTCAGT CTCCTCTTGT TITTATCGAGT ATTTATTCCT GTGCTTCAGT COGTAACAGC CCGAATTATC GGTGACCCAT CACTACATOG AGATGTTTOG ncerecense aathemeer caesteaath theastseme tithssenset eccentsith GIGCITIAGCA AAGIGGIGAA TGCCATITIGG TITICAGGATA TAGCIGACCI GGCATITIGAG GIATCAGGGA GGAAGCCTCA CCCATTCCCT AGTGTCAGCA AAATAATTGC TGACATGCTC ATCCATCTTG TCGGTCAGCT GGTTAGTCTC CTGCATATGT CCCTTCTCTA CTCACTGTAC SEQUENCE DESCRIPTION: SEQ ID NO: 124: ITCAACCITY TOCTGCAGGC TCTTTTCCTC ATTCAGGGAA SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1717 base pairs.
(B) TYPE: nucleic acid
(C) STRANDELNESS: double (2) INFORMATION FOR SEQ ID NO: 124: TOPOLOGY: linear 9 Œ 3 **æ** 2 2 2 22 8 33 6 5 20 55 8 2040 360 420 480 54 1620 1680 1740 1800 1860 1920 1980 2081 8 99 720 780 840 1320 1440 1500 1560 8 980 020 080 1140 1200 1260 1380 ATATOTCAAA TOTGAAACTG CTGTCTTTTA TATTAAAGTA ATTAAAGAAA ATGTATTGTG ATTGABATTA TTTTGACCTC CACAAGATGG CTCTATGAGT ATTCTTCCAG GGATTCTAAT AGGGCTTCGC GGTRACCAAG CGCTGCAAGG TGCGCCTCGT GCCGYTGCAG ATCCAGCTCA INCINCIAL ÁAATITATIT INATIAAATA CITRITAGAG GGITTIGAAA TGITTITINA CHACCTIGGG AAATCTTACA CCTTCAAGCA CTGTGTTTTT CTGCTGTGAT ATGCAGGAAA AAGGGCCCG GATTTTAGGA ATTCCTGTTA TTGTAACAGA ACAATACCCT AAAGGTCTTG GGAGCACGGT TCAAGAAATT GATTTAACAG GTGTAAAACT GGTACTTCCA AAGACCAAGT ATATICIDAA GCAAAAAGIC ICAGIAAIGA TITGGIAGIA ITAATITIGI GGICAITIGIT GOTTICAGACC AGCCATCAAG TATTITIGGGG ATATTAITAG CGTGGGACAG AGATTOTTGC ITITCAATGGT ATTACCAGAA GTAGAAGCGG CATTAGCAGA GATTCCCGGA GTCAGGAGTG ITICIATTAIT TGGAGTAGAA ACTCATGTGT GCATCCAACA AACTGCCCTG GAGCTAGTTG GCCGAGGAGT CGAGGTTCAC ATTGTTGCTG ATGCCACCTC ATCAAGAAGC ATGATGGACA SGATOTITICS CCTCGAGGOT CTCGCTCRAR CCGGGATCAT AGTGACCACG AGTGAGGCTG INCRECITICA GCIGGIAGCI GATAAGGACC ATCCAAAATT CAAGGAAATT CAGAATCTAA TIAAGCCGAG TGCTCCAGAG TCGGGTCTGC ITTCCAAAGT ATAGGACATT TGAAGAACTG CCOCTOCTGC TTACCTTCCT TITITIGITAA TGTGCTTTTA TITIATAAAA AAAATTACAA RGAAGATGCC TGTTTTGTCT CTACTGTGTA CTCTGATGGT ATCTTTCCAA AGTGCAGACT CTICIGAAGI ITICITAAAI IGIICACITI AAAGAAAAIG AGGIACCAAC AAIGAITIGG CITITADATI ACTODAAGAT GITATAANGI TAANGIGGAT GIAGIGCITT TACITITACAG ATTCATTGGA ATAAGATTAT TGCATATGAA TTTACCCACA GGACTCTGAA TCATGTTACC CACTCCCCTC ACANTOTIGT CCACTTAGIG AGTTGCAITIG ATCTAICGGT ACCAAAIGAT GIICAAIAAT IACAIAICIT ICIIGACIAI ACIGAITICI IAITIIGGIC ACIAIIIACIA AATCTCTGTT AATATTCTCT CTTTTAACTG AAAAGGGATG GGATAGAAGG GTTTGCAATG CCATATTATT GGTGGAGGGC TGTTTTAACA TCTTTGAAGT ATGGCTTGCT GAATATCTTT ACCAACATUT TGAATATATA TTCTAGTGTC CACAAGATTT AGCAAAAAGA TAAAGCTTGG GIGGAATATC ATTITIAAAAT GITCATGITC TOTICTAIAT ITTCITCACC TACTCTCCAA STATECTACT CACTESTEAA GEACASTCAS STEAAGGACT STAAGCCCAC ACAAGCTCTT CITATCICTA CTAGAATTAA AATGITAAGI CAAAAACGGC TCCTTTITTIG CGCCTCCTAG IGAAACITIAA CCAGCIAGAC CATITIGAGIA CCAGCAITITA GITACAAACG TCAAAGGCIT ATITIATITIAA GGINATAAAA TCITIGACAIT TATAATCITI C 345 8 2 2 2 25 3 35 6 5 S 55

1260 1320 1380

030 8 140 200

960

TGTTTGTGAG TCTCTTTCCC

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AGGACCACAG GITTITICIGC AGCIATITIC TAGCATITIC CAGICCCIGI GCCIGGACIG

1500 1440

COSCIDANCO GNACIOCCOT COCCATOTITI GOCIOCITICO TICOCOGRANG OCTOGIOCAN CCACGCGTCC GGTCACTATG TAGTGGAGGG GCAGACACCC TCCCGCAAAT TCTGGAAGGT TIGOTOCIGG TOACIOCOTT TATAGOCATT ACTOTOTIGE ATTITOTIGT TITICIGIGA AACACATACA TIGGATAIGG GAGGIAAAGG AGIGICCCAG CTACCACCCT TOGATGAATG GATTITIGTAA TICTAGCIGT TOTATITIGI GAATITIGITA ATTIGGAACAC TITIGITITITE TECCTIGIGCE ATTITACECTT CEACETTICE ATECTIGECT TOTTHISTOTE GACTHISGICA GTAGCCCCAG GACTCCTAGT CGCCGGCTTC AGGTCACTGC TITITOGICIC ICTIGCICCA CIGCAAAAAA AAAAAAA NINTITIGGN ANACATAATI TGAATAAAAT AATITITAAT GGATINTGNA AAAAAAAAA AGCCAACATC CTTTTOGAGC CATGAATATT GTCCGAACTC CATCTOTTGC TCAGATTGGA OTCACGAATG GGAAGCCAAG TGCCATCTTC AAAATTTCAG GTCTTAAATC TGGAGAAGGA TOTOTOTACT TITOTIATOC TGATTCAAAT GGAATGCCAG TATGGCAACT COTAGGATTT ACAGCTOCAC AGCAAGTOOC AGAGGATAAA TITOTITITO ACTIACCTGA TIATGAAAGT (2) INFORMATION FOR SEQ ID NO: 125: COGGCAAATG TGGTTCTGAA ATGGTATGAA AACTTTCAAA GACGACTAGC ACAGAACCCT OCTICATICAT TIGGIGICIC ICAGGCCCAG ANGACACCAA GCCCANCIGA AANGIICATI TOCTOLOGITO ACTOLITICAC TOLOGITCACA CAAAAGAIGI TOGACAATIT CIACAATITI ATCAACCATG TIGIGGTTTT TATGCTGGGA ACAATCCCAT TICCTGAGGG AATGGGAGGA ATTICAGIGG AATTATTAGA CAGTAIGGCT CAGCAGACIC CIGIAGGIAA IGCIGCIGIA (1) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 125: (A) LENGTH: 431 base pairs TOPOLOGY: linear STRANDEDNESS: double TYPE: nucleic acid 126: TICTIGIAAC TCAGGITAGG 1680 1560 1717 1620 120 540 480 420 360 8 240 180 720 660 90 804 780 25 6 35 ಜ 20 8 ន 8 45 2 5 GOGTTATTTT G GOCCATOGAG GCATTICITY CAGOGAAATG OTCCATNATY TCAGCCAGAA GCCATTOCAT GIGGGCAGGI CAGIGATIGIC AGCAGIGGAG TGATICCCAG CACAGCGGCI ICIGGGAAGA GCTGGGGTQA ACTITATITT AGCCCTTCCC TIGTTGCTCT TATGGAAGAA CAGAGGAGGG GRAGOGICIO GRATOGOGCI OCCCCIGANG OCCCIGANOT GRAGIACCIT OCCAGCATCI GOCACAGECE AGGGEETTGA AGCEAGETGG CECTGGAGAG GGGETGETGT GECAGETTGG THIGHTACCA CAATTAGAAC TOOCAGAGCA CTGAAAGAAA AGACTITTOCT TCCCGAAGAT CTICAAGGAG GGGGCAAAAN GACCITIAAG TITTITAGGIT TAACACAGGG AACCCINCAAA TANGTTANOT CONSCRIPT TOTOGCCCAG CTCTOTOTTA TINAGGSCCC TTGSCGAAGA CCAGAAGGAA CTOGNGAAAAG CGAACAGGAA CCGTTCAGGT CCTGAAGCGG TCAGGCCGAG GATGCTTTGC AGCCATTGGT TGACTGGTTA TACAAGGTGG AGCCACAGCT GGCTGAGGAC AMPICTOTOG AGCGGCAGCA CAAGTTGGAG GAAGCCCTGC TCTTTTCGGG TCAGTTCATG ASTCAGAAAC TIGACAATIT CCIAGGAGAA GICAGAGACA AAIGGGATAC IGITIGIGG AAACTTCAGC TTTCTAAGCA TAAGGAGTTT CAGAAGACTC TTGGTGGCAA GCAGCCTGTG TCTCTCATTG ACACCCATAA GGAATTCATG AAGAAAGTAG AAGAAAAGCG AGTGGACGTT GCACTOGOTO GGACACTOTO TOTAAACTOT CTGTTTOCAA ACAAAGOOGG CTTGAGCAGG CAGCCCCGTGC NGGCACGAGG AGAGTCACCT GGACTCAGAA CTAGAGATAT CCAATGACCC AGACAAAATT 5 AMBCADAGCA AACGCTTOGC TTTCGGGDAG CACTTCCTOG ATBACACAGA GGCCCTGCAG CCTTAAAACA AGCGGAAGTG TTTCGAGACA CAGTCCACAT GCTGTTVGAG AGCTGATTGA GAATAGTCGA GATGACACCA CTTGGGTAAA AGGACAGCTC CAGGAACTGA INFORMATION FOR SEQ ID NO: 127: Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 126 X. SEQUENCE CHARACTERISTICS: ACCOCICIACC TIGACCICCT CATGAACCIC AIGGAIGCAC ACAAGGITTT SEQUENCE DESCRIPTION: SEQ ID NO: 127: (C) STRANDEINESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 3752 base pairs TOCCTTICE

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INFORMATION FOR SEQ ID NO:

660 600 540 480 420 360 300 240 180 120

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WO 98/39448 348

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

120

180

PCT/US98/04493

₹	WO 98/39448	PCT/US98/04493	*	WO 98/39448	PCT/US98/04493
	349			350	
	AACTCAGCAG TAGCCATGGG AGAAGTCATC CTGGCTGTCT GCCACCOCCA TTGCATCACA	840		ACTICIANAS ANTANTICIS TINIGAAGCT OCCTINITIT TITITCTITIT GPAAGTING	2640
,	ACCATCAMAC ACTIGATICAC CATCATCCGA GCTCGCTTCG AGAIGGTCCT CACATGGGCT	006		ATTITICATOT GARIATITAL GIAGATAAA TITOCCICCT GGIAACCCTG TAATGGAIGG	2700
^	AAGCAGCACC AGCAGCGTCT TGAAACGGCC TTGTCAGAAC TGGTGGCTAA TGCTGAGCTC	096	ν,	GCCCCAGNA TGANTATT GAGAAAACA AGTGAAAAGG TCAAGATACA AATOTGTATT	2760
	CTGGAAGAAC TTCTGGCATG GATCCAGTGG GCTGAGACCA CCCTCATTCA GCGGGATCAG	1020		AAAAAAAAA AAGCTATTA ATAGGGTTC TGCGCGGTGC AGGGTTGTAA ACCTGCTTTA	2820
0	GAGCCAATCC COCAGAACAT TGACCGAGTT AAAGCCCTTA TCGCTGAGCA TCAGACATTT	1080	01	TCTTTTAGGA TTATTCCTAA ATGCATCTTC TTTATAAACT TGACTTGCTA TCTCAGGAAG	2880
	ATGGAGGAGA TGACTCGCAA ACAGCCTGAC GTGGACCGGG TCACCAAGAC ATACAAAAGG	1140		AIRAATTATA TTAAAAAAT AAGAATCCTG CAGTGTTAA GGAACTCTTT TTTTGTAAAT	2940
4	ANAMACATAG ACCCTACTCA CCCCCTTTC ATAGAGANAT CCCCCAGGGG AGGCAGGAAA	1200		CACGGACACC TCANTIAGCA AGAACTGAGG GGAGGGCTTT TTCCATTGTT TAATGTTTTG	3000
2	TECCTAAGTC AGCCAACCCC TECTECCATG CCAATCCTTT CACAGTCTGA AGCAAAAAC	1260	15	TGATTITING CTAAGAGAG GGAACCTCAT CTAAGTAACA TITIGCACATG ATACAGCAAA	3060
	CCACOGATICA ACCAGCITITE TOCCOCCTGG CAGCAGGTOT GOCTOTIVAC ACTIGGAGGGG	1320		aggaoticat tocalaacto ictitiggata tiotiticaot actososott taaaggacaa	3120
22	CAAAGGAAAC TGAATGATGC CTTGGATGGG CTGGAGGAGT TGAAAGAATT TGCCAACTTT	1380	20	atroctocta gaattcagge graatgtaa gtottcagaa aaggtcagaa catttggget	3180
	GACTITICATIG TCTGGGGGGA AAAGTATATG CGTTGGATGA ATCACAAAAA GTCTCGAGTG	1440		TITAAACIGA TITOTTOCTC CCTATCCAGC CTAGACACCA GTAACTCTTG TGTTCACCAG	3240
, v	ATGGATTTCT TCCGGGGAT TGATAAGGAC CAGGATGGGA AGATAACACG TCAGGAGTTT	1500	1	GACCCAGACC CITGGCAAGG GATAGGCTCG ITGGTGACAT TOTGAATITC AGAITIGITT	3300
3	ATCGATGGCA TITITAGCATC CAAGITCCCC ACCACCAAGT TAGAGATGAC TGCTGTGGCT	1560	25	INICCACITI TITIGCIAIT TAITIAAAIG GIOGAICAAC TICCCACAAA CIGAGGAATG	3360
	GACATITICG ACCEAGAIGG GEATGGTIAC ATTGATTATT ATGAATTICT GGCTGCTCTF	1620		ANTICCAGGA GCCTGTTCTG AAANTGTGGA GGTAAGAGAA ACAGGTGCTC GTCCTTTAAT	3420
8	CATECEAACA AGGATGOOTA TEGACEAACA AECGATGEAG ATAAAATEGA AGATGAGOTT	1680	30	GGAOTTCACC AGACACTTG TTAACCAGTC CTOTTTAGCTT TOGTCTTTTT TTGTGGGTAA	3480
	ACAAGACAAG 16GCTCAGTG CAAATGTGCA AAAAGGTTTC AGGTGGAGCA GATCGGAGAG	1740		TAAAGTCAAC TGACCAAGTG ACCATGAAAA GGGGCTGTCT GGGGCTCCTG TTTTTTAGCT	3540
v	AATAAATACC GGTTCTTCCT CGGCAATCAG TTTGGGGATT CTCAGCAGTT GCGGCTGGTC	1800	;	GETOTICITIC AGCICCIACO ATOTICCIOT OTGATIATOT CAATTGOTIT TAATTGAGGO	3600
ž	COTATTCTOC OCAACCOTGA TGGTTCGCGT TGGTGGAGGA TGGATGACCT TGGATGAATT	1860	35	agalactgaa gcyctaccaa tgaactgyty agalacaaga cacactyyyg taftaalaty	996
	ITTAGIGAAA AATGATCCCT GCCGAGCACG AGGTAGAACT AACATTGAAC TTAGAGAAA	1920		GCTTGCAGTA ACAAAAAAA AAAAAAAAA AAAAAAAAA AAACTTGAGG GGGGCCGGT	3720
9	ATTCATCCTA CCAGAGGAAG CATCCCAGGG AATGACCCCC TTCCGCTCAC GGGGTCGAAG	1980	40	ACCCAATICG CCSTATATGA TOSTAACAA TC	3752
	STOCARAGEA TETTECESSS CAGETTCECE TACTOSTICE ASCITCEASTS CTASTICAGAS	2040			·· · <u></u>
4	TAACCACAGC TGTACATCCA TGCCATCTTC TCCAGCCACC CCAGCCAGTG GAACCAAGGT	2100			
5 .	TATECCATEA TEAGGIAGEA AGTIGAAGS ACEAACACA ACTITICAIT CTAGTEGGAC	2160	45	(2) information for SEQ id no: 128:	
	ATCCTTGCT GOTGATACCA GCAATNAGTT CTTCCCCGGC CTCCACAGGT GCCAAAACTA	2220		(1) SEQUENCE CHARACTERISTICS: (A) LENOTH: 1144 base pairs	
8	ATCGGGGGAA CCTAAAAAG TCTGCGAGTC GCCCTGGGAG TCGGGCTGGG AGTCGAGCCC	2280	50	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	GENETICENSC CACCACCCCG CANCANAGTS ACSCITICTGA CITTGACCTC TTAGAGACGC	2340		(D) TOPOLOGY: linear	_
	ATTECTTOTT CCGACACTTC AGAAAGCAGC GCTGCAAGGG GCCAAGGCAA CTCCAAGAGA	2400		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 128:	
22	GGGCTAAACA AACCTTCCAA AATCCCAACC ATGTCTAAGA AGACCACAC TGCCTCCCC	2460	55	TRACCCICTO CCTGCCGGGC TCAGTGCTGG ACGCTTTCTG TITTTGTCGCA GTCGGTCCTC	09
	AGGACTICCAG GTCCCAAGGG ATTAACACTGT CTAAGGACCC CCAAGCCACT ATCCACTTTG	2520		GGINACACCA GCGGCCTGTG GTCCACCACT CCATTCAGCA GCTCCATTTG GTCCAGCAAC	120
8		2580	9	CTINGCAGGS CCTTCCCTTC ACACTCCAG CAACAGGCT GACAAGCATC GACCTCATGG	180
			;		_

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35 23 50 3 8 3 20 5 8 55 0 CITAGCITICC AGGITICACCC TAACCCIGIA CTAACCIGCT TGGTGGACTT GGAAAAGACT TACATTICAA AGAAAGGTAT GTIGICTAAC AGGGGACCAA CAGAAGGTAG TATTGACAAC GITGITGGIG TIGICAAAGI TIGITTITIC TAAAGITGIG TGATTATATA TITIGACATIT GAGATTTIOCT CTGACTTTAT TTATATGGCA TGAAATCTCT GGTTTATTTT GGGATTTTTT AGCTGATGGG AGCAGCTGGT GCCTGGCCTT CGGCTCCTGC GTCCCCAGAA CCCAAGGGAA 5 CGITAAAITC TIGATAGAIT TITTATTAIT GOGATIGGCCC AGGOCCAGCG CATTOTIGCAC TOGTTTACTT TAAAATGTAC AGATTCTTCT TCACTOTATC TIGGATCAGA GACAAAGGAG GACCCGCTTT ACCCCTGCTG CGGGAAATGG CHOCHROCCC CAGGOCHOGG ACCHIGAAGC CCHCCCGGCA ICHOGCATICC GAGCCTCCCG GOGAGGAAGC GGGOGGACGC CTGTCACCCC TGGCAGGTGG TGAGTTCAGG TGGGGGGCTCC GACATCACCA GCCAGAGCAC AGGAAGCCAC CCTGCCTGCT GGGGAGGAGG GACCCACACA COGGCOOTOG CTICIOTGAT TIATITICIT GATOGIAACT TCICAGAGCA GGGCRATIGG GOGGATOGAG CCCCACCTGA GGTGCCGTGT CACACGGGTT AGAGGGTCAC TGGGAAAACAC CICICITICA GIGACIOSCI ACAAGOSCCI GAGAGOTIGG CAGCCAGOTI TOGAGCIOGA TOGACTOTOA TICCITCCCC CACCCICCAI ICICCAGCGG TIGGCCGGIG ITAGAACICG AAAAAATGIC ATIGITICCT GITTGITAAT ATTAGGGTTG TAAGGTGICG TITTGAGGTA TGTTCCTGCT TCTACTAAAA AAAAAAGAGC ACAAAAGAAA AACTAAATTA TTGAAAAAATT AGAGECACAE CAGETETIGGA CATEACEGEE CETIGGAACTIG GGGCCAECAG CEETIGGGCAE TOUCTETOTE GOGAMAGGAG AGACGGGGCC TECATEACGC CTGTTACEAG AGGATECECG AACCCCOOTT TCCTTCAACO TCCACATTCC AGGTGACCAC ACGTGTCTCC TCCTCCTCAT COTCATOGAG GCCACATOGG GCCACCCOGC TCCCTCGGGA TOGCTCCGCT GCACTTITTGA CTGGGCCGGC CCATTCTTCA CACGCCTGCC AGAAGCTGGA GGGGTGCTGG AGACCCATAG COCTOCAGGG TOCOCTTOCCC TOTOTTOCCG CAGCATACAC GAGGGCAGGC AGTGGCOTTO AGECCECTEG GEAGTITICTE CECCEAGETT COGTATGCET TEAGGGAAAAG GTEACAGETG INFORMATION FOR SEQ ID NO: Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1864 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 130: 1260 1200 1140 1080 1020 1830 1800 1740 1680 1620 1560 1500 1440 1380 1320 960 900 840 780 720 660 600 540

8 S GETGGGEEGE GGCETETGGA GETGGGATTT GGGAGGACAC AGCAGGCAGE GETGGEETTE ACGGGTTGCT GCCCAAGTGC ATCATGCAGG CCACGGACAT CATGCGGAAC AGGGCCCAAG GCATOCAGAG GAGCACCCTG AGCGTOTYCC TGGAGCAGGC GGCCATSTTG GCACGGAGGC TOCCTCCTTC COGGACAAGC CTGGCCACCC TCGCTGTGAT GACGAGCTGG CTGATTGGCC TOCAGGGATG GOCCAANGOT TOCGGAROOG COCGTTOCGG GACCTGCCCA GOGTOCTOCC CCTCTACCOC CTCTGCCAGC CGCCGGTOGA TGGGGACCTC TGAACACCCA AATGCCCCAC OGTIGGAGATT CTIGGCCAAAA ACCTIGCGAGT CAAGGACCAG ATIGCCCCAGG GTIGCTCCGCG INFORMATION FOR SEQ ID NO: 129: $\widehat{\mathbb{E}}$ Œ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 129: (A) LENGTH: 1830 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDMESS: double
(D) TOPOLOGY: linear 90 240 180 120 420 360 6

30 CAAT agatacaaga tigaatotot atticitaaa aatacaacti tototiotac titgaaataa ATGATOCTYY TYTCAAAAAA AAAAAAAAAA AAAAAAAAC TCOAGGGGGG GCCCGGTACC

25 8 5 TCTARTITTA AAACTGTATG AGGACTTTGT OCTGAAAATA GAGTATTTTT TTAAAGTAAG ACAATTTICT TAAAITACCA AGITIGGITT TIAATAATIT CICAATAITA TOCGCCAAGA CITICCCOIT ACIGCOTTIT CACCACCIOT CITICCCCATO CITTATTIAT CIGIATGAAC CTTTTAGAGA AAAGICTGGA CTCAGCCACA AACICTAATA AGACCTGTAC ATCTGAGAAC AACATTIGIA TACCIGAACT TATTITAAAG ATGAACIGAA ATGCACATAG CCAAGTCTIG TGTAAAAACA TICAGTYGAG ACCATATGCA TITTCTGTGC TGTTTGTACT TGAGGTATGT OCTOTOTOG TTTAAAAGCA GATTACAGAA ATGTAAGTCA ACTTAAGAAC RGTGAATGAA ACAGATITIGA CATTACAGCT AAGGAAATAA TITGAGTTGA TICAGAAATC CTGGCATGTG 1144 1140 1080 1020 960 900 840 600

780 720

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GACCACTOGO TICTITICAGA TOTITOTIGO AATTATGATA ACATGAGATI TOCTOTIGIG

540 480 420 360 300

TOCAGOTOCO CTOCOCACCO TOCTOTTOTT TOCAMACAG ACCOMAGOAG GOCAGGOTOM

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GIGGOCCCIC GICTAGATCA TGAIGIGCCA GITICIGAGA CAICIITITA AGGCICITAC

CTTOGTCTAA TTCGCACTTT CCTCACGAGA ATTAAATTAA GCAAAAAACA AACAAACATA

S

GACAGACCTA CAACCCCTIGG CGGATATGGA GCCCCACGAT TGGAAGAAGA AGCTCGGACC GCACAGAAAA CTCCCCTGCT CCTCACGCTC CCTCCACCTC CAGTCCAGCT GACGACTTGG

PCT/US98/04493

Per 198/39448	PCT/US98/04493	WO 98/39448
353		354
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 130:		TCATTICCAG GGGTGGGAAT TITITITIAA TANGTOTCAR GAATAAAGIT GITTITIGAAA
GOCCOCCOGO ATGOCOACC CAGCOTOGGC CCCAGACACA CGGGCTCTGG TGGCAGACTT	9	akaaaaaaa aaaaaaaa aaaaaaaa aaaaaaaa aaaaa
TOTAGGITAT AAGCTGAGGC AGAAGGGITA TOTCTGTGGA GCTGGCCCCG GGGAGGGCCC	120 5	5 AAAA
AGCAGCTGAC COSCTGCACC AAGCCATGCS GGCAGCTGGA GATGAGTTGS AGACCCGCTT	180	
COGCOCACC TICTCTGATC TGGGGGTCA GCTGCATGTG ACCCCAGGCT CAGCCCAACA	240	Ś
ACCTITCACE CAGGICTICG ATGACTITIT TCAAGGGGGC CCCAACTGGG GCCGCCTTGT	300	JANT (7)
ACCEPTETIT STETITIOGGE CICCACTOTO TOCTGAGACT STCAACAAGG AGATGGAACC	360	
ACTIGOTOGGA CHAGTIGCAGG AGTIGGATGGT GOCCTACCTG GAGACGCGC TGGCTGACTG	420 15	<u>e</u> 0 (
CATCACACC AGTECROCCT GATTATCCCA CATCACTGAA GCTCAGATGG CTCATGAAGT	. 480	(b) YOPOLOSY: Linear
AATTIGCAGT GAAATTITAA GCGACTGTGA CITCIGCTGCA AGTICCCCAG AICTIGAGGA	540	(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 131:
CONCERNOT SITESABATOTIC CARTICAGES CATTESAGGAS GAAGCTICAGA ACCTIDAAGGA	20	GOCACGAGGG CGCGGCAGGG CCCTGGACCC GCGCGCTCC CGGGGATGGT GAGCAAGGCG
		CTGCTGCGCC TCGTGTCTGC CGTCAACGGC AGGAGGATGA AGCTGCTGCT GGGCATCGCC
GCIACAGAAC GAGGIAGAGA AGAGATGAA TATGAGICCA CCICCAGAGA ATGCIGACCC		TIGCTGGCCT ACSTCGCCTC TOTTTGGGGC AACTTCGTTA ATATGAGGTC TATCCAGGAA
GOTCATICATIC TCCATTGAGG AGAAGATGGA GOCTGATGCC COTTCCATICT ATOTTGGCAA	720	ANTIGGIGAAC TAAAAATTGA AAGCAAGATT GAAGAGATGG TTGAACCACT AAGAGAGAAA
TOTOGACTAT GOTOCAACAG CAGAAGACT GOAAGCTCAC TITCATGGCT GTGGTTCAGT	780	ATCACACATE TALABABAC CITITIACCAC ARABACCACACA ARABARCACA
CARCOSTOTT ACCATACTOT GTGACAAATT TAGTGGCCAT CCCAAAGGST TTGGSTATAT	840	
AGAGITICICA GACAAAGAGI CAGIGAGAC ITICCITIGGC ITIAGAIGAGI CCCIANITIAG	006	ANGENTOGEA AANGAATITIT GATAACAGGA GOOGCAGGGT TOOTGGGGCTO
AGGAAGGCA ATCAAGGTGA TCCCAAAAGG AACCAACAGA CCAGGCATCA GCACAACAGA	096	GACANACTICA TGATGGACGG CCACGAGGTG ACCGTGGTGG ACANTITICIT CACGGGCAGG
CCGGGGTTTT CCACGAGCCC GCTACCGCCC CCGGACCACC AACTACAACA GCTCCGCTC	1020 35	AAGAGAAACG TGGAGCACTG GATOGGACAT GAGAACTTCG AGTTGATTAA CCACGAGGTG
TCGATTCTAC AGTGGTTTTA ACACCAGGCC CCGGGGTCGC GTCTACAGGG GCCGGGCTAG	1080	TOGRACOCCT CTACATCGAG GTTGACCAGA TATACCATCT GGCATCTCCA GCCTCCCCTC
ACCERCANCE COMMENCE CITE CONTRACTAL ABACTICITAL TRACCACCION ACAGEMICAA	1140	CAAACTACAT GTATAATCCT ATCAAGACAT TAAAGAACCAA TAGGATTGGG AGATTAAAGA
	40	TGTTGGGGCT GGCAAAAGA GTCGGTGCCC GTCTGCTCCT GGCTTCCACA TCGGAGGTGT
AAAAGAAGAA AGAAGAAAAA AAAAAAAAA I IAAAAAAAA	00 00 00 00 00 00 00 00 00 00 00 00 00	ATGGAGATCC TGAAGTCCAC CCTCAAAGTG AGGATTACTG GGGCCACGTG AATCCAATAG
MICHIGALIO ANAMANATA TITITIANA ANAMANIATA CITICANIA GOGGEGIA		GACCTOSGEC CTOCTACGAT GAAGCAAAC GTOTTGCAGA GACCATOTGC TATGCCTACA
CCATAACTAA CTGCTGAGGA GGGACTGCT TTGGGGAGTA GGGAAGGCC CAGGGARTGG	1320	TGARGCAGGA AGGCOTGGAA GTGCGAGTGG CCAGAATCTT CAACACCTTT GGGCCACGCA
OGCAGOGOGO TOCTTAITCA CTCTGOGGAT TOGCCATOGA CACOTCTCAA CTGOGCAACT	1380	GCAGGGGTC
GCTTGCCCAT GTTTCCCTGC CCCACCCCAC CCCTCTTCTC CGGCTCCCTG CCCCTCCAGA	1440	
TIGCCIGGIG ALCIAITING TITICCTITING TGITICITIT TCTGITITICA GIGICITICE	1500	
THECAGGITT CIGINGCCGG ANGAICTCCG TTCCGCTCCC AGGGGCTCCA GIGTAANTTC	1560	MANTGOCCT COTOGOTOTO ATGACAGOA ACOTOAGOAG COCOGOTOAAC CTGGGGAACO
COCHICOCO RESESABATIG CACTACOTTG TITTIGGGGGG TITTAGGGGGTG TITTIGITIT	1620 55	CAGAAGAACA CACAATCCTA GAATTTOCTC AGTTAATTAA AAACCTTGTT GGTAGGGGAA
SUMMERCALLY CONTRACTOR TO THE STATE OF THE S	1680	STRAMITCA STITCICTICS GAROCCEAGS ATGACCACA GARAGABA CCAGACATCA
Contraction of the contract of		AAAAAGCAAA GCTGATGCTG GGGTGGGAGC CCGTGGTCCC GCTGGAGGAA GGTTTAAACA
ACCORATOCO ACCAACTOCO AACAGCGAGG TOCAAAUSTOU ATTITUTTA TITTITAAC	09	AAGCAAITCA CIACITCCGT AAAGAACTCG AGTACCAGGC AAATAATCAG TACAICCCCA

CACTITITICA ATCICTCTT TIATOTAAAA TAGCOTAGAT GCATCTCTGC GTATTTTCAA GCTTTATGIT TCTCTTTTAA TICAGAGTTT TICCAAGGTC TACTTTTGAG TIGCAAACIT GITTITITAT CITICCIOTGA GAGCATATOI TOTGACTOIC GITGACAGIT TIATITACIG TOCTITAATG AAATGGATGT GCCTAAAAGC TCCCCTCAAA AAACTGCAGA TTTTGCCTTG GTTTAAAGAA AGACTITAAC AGGTGTCATG AAGAACAAAC TGGAATTTCA TTCTGAAGCT GACCOGAGCT GOGAACOGGA ATGGCACAAC CAGTCTGAGA AGGACAGTTA TGGTTACATT CTOCCTATCT GOOGCOOGG CAGOTTOGGG GGCACAAAGT TAACATATTC TTOGTTAACC GACTITGAAA TATICCIGIT GGICATGAIC AAGGATATIT GAAATCACTA CIGIGITITG TIGICAGGIG GIGGIGGGC GGCATIGATI TIAGGGCAGA TAAAAGAATI CIGIGIGAGA AAAAGTOOGT ACTTAATAAA TGAGTOGTTA TACTATOCAT AAAGAAAAAT CCTAGCAGTA GITICITIGI GAAGCIGAAA AGGAACAITA AGCGGGACAA AAAAIGCCGA TIITAITIAT AGGACACAAG ACTACCATTG TACACTTGAT GGGATGTATT TYTIGGCTTTT TTTTIGTTGTC AACCAAAGCC TOCCAGAATA AAGAAAGGAC GGACTCOCCA CAGCTGAACT CCTCACTTTF TACACTGATG GTACATTCAG GNTCCCTCGG CCAAGGACTG GACCAGAAGA ACACTTGGGA TICCIGAGCA ACAAGGAIGG GCICCIGGGI ICCAGAIAACA AGAAAGCIGI AITCAGGGAA TACCARGOTG CAAGAATCTA CTATATCATG GCAGAAGAAG TAGAGTGGGA CTATTGCCCT 2 CCACTOOCTO CTGAGCCTGG TGAGGTGGTC ACTTATCAGT GGAACATCCC AGAGAGGTCT ATCTTOGGTC CACTTATCAA AGGTGAAGTT GGTGATATCC TGACTGTGGT ATTCAAGAAT GOCCCIGGOC CAAIGACICT OCTIGIOTIT CCIGGAICTA TIATICIOCA GIGGAICCCA AATGECAGEE GEECETACTE TGTGEATGET CATGGAGTGE TAGAATETAE TACTGTETGG TCAAGGACAT GTATAGTGGC CTGGTGGGGC CCTTGGCTAT CTGCCAAAAG GGCATCCTGG INFORMATION FOR SEQ ID NO: 132: Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 132: 909 (A) LENGTH: 2012 base pairs TOPOLOGY: linear TYPE: nucleic acid STRANDEDNESS: double 2040 1980 1920 1860 1800 1740 1680 1440 2041 1620 1560 1500 240 540 480 420 360 300 180 120 6

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6 ႘ မ 25 8 8 3 2 5 GAAGAAGOCA ATOTGAAGAT GCTGGGCATG CAGATCCCCA TAAAGAATGT TGAGATGCTG TTCCTCTATC GGAATGGCGA GAACTACCGG GCAGATGTGG TGGATCTGTT CCCAGGGACT TACATECTOG CCATEGGCCA AGATETOGAT CTACACACCA TCCACTITICA TOCAGAGAGC GOGAAACTOT ATGCCAACCT TAGGGGTOTT ACCATGTACC AAGGAGAACG AGTGGCCTGG GOCAGTATTA ACCTACAGGA TGAAACTTTC TTGGAGAGCA ATAAAATGCA TGCAATCAAT GATGAAAATA AGTCTTGGTA TITIGGAGGAA AATGTGGGCAA CCCATGGGTC CCAGGATCCA NAGCCCCATG GAGGACGGAN TGACATGGAT CGGGAATTTIG CATTGTTGTT CTTGATTTTT TATATTTCCT TCTGACACTT GGAAGGTATT GAAATTTCTA GAAATGTATC CTTCTCACAA GOGGCATOGG TOGTGGAGAA GCAGAAGGAG CAATCAAGCT TATCTGGATA TTTCTTTCTT GACAGOTTICA AGOTTICTIGTO TITICAAACAG TAACATICTIGG AGOOTIGGAGA TATOOTICAGG GCCICIOTIT IOOTIGCCAT IAGIOICACC CITCIOCICG TIOTICIOGC ICTIOGIGGA TTAAGCCCTC TCACCGTCAT CACCAAAGAG ACTGAAAAAG CAGTGCCCCC CAGAGACATT ACTGACCATG TOCATGCTGG CATGGAGACC CTCTTCACTG TTTTTTCTCG AACAGAACAC TITIGAGGITG TOGAGATIGGT GGCCAGCAAC CCTGGGACAT GGCTGATGCA CTGCCATGTG TAAGAATATA GOCTTGATGG GAAATTGAAG GTAGGCTGAG TATTGGGAAT CCAAATTGAA ATTTCACTIT GAACTGAGGC CAAGTGAGCT GTTAAGATAA CCCACACTTA AACTAAAGGC TATTTATTTT ACATOGAAAT AATATGATTT CACTTTTTCT TTAGTTTCTT TOCTCTACOT AAGCACATOT GTAGTGCACT COCAGCAGGC CATGGACTAG TCACTAACCC CACACTCAAA GTOGTTTGGT ACCAACATCG ACAGAGAAAG CTACGACGCA ATAGGAGGTC CATCCTGGAT AACCTOTOGO ACTGAAAGGA ATOTTGAGTT ACCTCTTCAT GTTTTAGACA GCAAACCCTA TGAGCATGTA CAACCTCTGG AGCTAGAAGC TCCTCAGGAA AGCCAGTTCT CCAAGTTCTT GGGCACCTOG CACTAAGGGA GTACCTTATT ATCCTACATC GCAAATTTCA ACAGCTACAT TCCATTAAAG TACTTGTTAG AACACTGAAA AA TITIGATICE ACTAGAGACC AAGAGAAAAA CTCAFTGATT GGGTTTCTAC TTCTTTCAAG GACTCAGGAA CCTTGGCAGT GAACTACTTT GAAGAAGTGG TCAATGGGTT GTTGCTGCCA 1980 1740 1680 1620 1560 1500 1440 1380 1320 1260 1200 1140 1080 1920 1860 1800 1020 2012 600 80 840 780 720 660

Ε SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 133:

(A) LENGTH: 1669 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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WO 98/39448

PCT/US98/04493
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(X1) SEPUENCE DESCRIPTION: SEQ ID NO: 133:	
GAGCAGTATT TTAACCAACT TOTATTACAG ATOTTACAGT TCATOTTAGG AAGTCAGAAA	09
AGACITIOTI TOTCITIOTI CIGCIGATOT GAGICATOTI TIVITOGOGIC TICCATGGA	120
CATTIACCTG TTGCTCCGTC CAGATGTTGA GGGCCAGTCT AGGCTGACAC ATCCTACCCG	180
AGGACAAGCC TGTTCTCCAT TTCTTCACTC TCCCCTCCCC	240
THAGATTACC STITTCGACG ACAGATTAAC CAAAAATGCC CCACACAGGT TTTATTACTG	300
TTATATACTA TACTITITAAC AGIACAGACC CTAAAITTTA TTATITIGITG CTCCCCCAAT	360
CTGATACCAA ATGTTTAAAG TTGTTTGAAA TCCAAACATG GTAGTGTTCA 1GGGTAAATA	420
TITICINGGC TATGTAAGAG TTAGCAGCCC ATAGCATAGA AGTAATCAAG TAGCATCTGA	480
GACTIGITIGGA GCCACTAGGG CCTCTGGG CCTAACAGCC TCACTTCCCC AGCCTCACCT	540
TOCTISTICCTC TOACACTISCS ATCAGGGCTO TTAGTIGGCAC CTGTATGAGG CCAAGTISTGC	009
GTCCAGGGGA ACAGCACAGG TTAATGGGTC TCCCTAGAAC TCATGAAGTC AGTTTAATTC	099
ATGCATGAAC ATGAGITCAI TITAIGITIT ATATAGCITT CITAGACATA CCAAACCATC	720
AFFCATAAAT CAGATAAATT AFFCAGTTTT TOTOTTTAGA AAGCTAAGTA TOTOTAGCTG	780
GANACAAAAA TGAGCOTGTT TTCTCTCCTG TTAATCTAGA GTOTGCAGTT ACACATOST	840
GGATAATTIC ATGITCCAGG GGCGCTTGGC ATCITCCCATG GACTGATTCC CAGGAAGAAA	006
AGCCCAAAGG GAAACCCACG ATTCCTTTCG AGTAGATGTG GGAAAGACC CATTGGAGGA	096
TATCAGGICC TOTCANAITC AGITOTOTOT GTGGCTCCTT GTTAGCAGTC ATOTTCACAL	1020
GOTOTINGGA GOCTCCCCAT CCACCCTTA CATCATOTAG GOACCAGTOT CTTOTGAGAT	1080
тамссттово асасмотово ттасстова ваамтемва овосствест бвассемово	1140
AGAGGAGCCA GTGACACAGG CAGAGGGGTG CAGGCCTCCT TCCCTTCCAT TTGGAGGAGG	1200
TGGTGCCAGG AGCCTGCCC CTTACCTCTG CTGAAGCATA AGTGGACTTT GCTTTTTGGG	1260
CTINICICIO AINCAIGCIG GAGCCCIGCC ICTCCACIGC TAGAIGGAAC CIGGAAICTC	1320
TCATICTACCT CTTAGTCTGT CAGTTTCTAC GTGTGAGAAG CAAGCTTGTG GGCCAGTGTC	1380
CTIGIACATO CTOTAGCACT TAAAAATAA TTCCAGGGTT CCCTGGAAAA CCAGTCCCAG	1440
GGITCCTATG ATCTGTAGIT TCTACCTGGA TTATAACTGG TTTTGGGTAC CTGAATTTTG	1500
ATTGGTTAGC CTTAATTATA GTCTGGCGTG ATCATGTAGA ATCTTTTCTG GTGAAGAGAT	1560
CATAAAGTTC TATCAAGGAG TTCTATCAAG GCATCCATGT CAGTGGTGCT ATGCTGGTTA	1620
CAACTIGAGA ITITIGAAAT AAAAAAITIG ICAFAAAAAA AAAAAAAA	1669

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CACTITITISCI AIAIAACCIA AGIGAIAACC CICITITIAGF IACCIGCCAA ACTOIGANCI TGGITTATAT TGCAGITAAC ACAGITACAA AGCIGIAATG GIGICITITI INCCITIIGIA ACGGRATGIG TAAATCAAAG TATATACATT GIGIGGIGTT CCTGITICIG GAGITICAIG AGGATITIACA CATGGCATIC AGTGTTCTGT ATAGATCTGC CTACCTTTGT GAATTCATCT TIGGAAAAGI TGAGAACIGC ITICICIACC TITIGGAAAA TAATIGATAI ICCATAITIGG GITAACCCCT CITCCITIGA GAGACACCG GCGATGGIGG ITAACTCCTT GICITITICTC TCTCTCCTAC TGGTTATTCT TGAATTAAGC ACAGACTCGT CAGCTCGGTT GCTTTATCAT GAATAATGTG TGTGACCTTG CAGTTCTTCC ACAGTTCAGC AAACAAGTGC TAGGTTCACT GACCAMAMAT TAAGGAAGGA AAACACAGTT TTTAAAAGGA TCCATCTTTT AACAGGCGAA CATTITITIAA TACTCIGAIT CIGIAACAIT TCIGAGITIT GITTIGITIT ACAGNAAAA ACCGATICTION CTATIGGTIGGT GCACCTTGCT GTTGTACTTC TGAAATCAGA COTGTGTGAA CGATCATTIC TGACTTAACC GTGAGATGCT CACGAGTACC CTTCCTGTTG TITTIGTTAGC AFTGAAATCG AGACTATTTA TFTGGAATAT ATACAACAGT GTTTTTCCAC TGTATTTCAT ATTOTCADAG ACTTOGATAT GGIGAACCTA TTAAACCTAG AAATTGTATT CATCCTTTCA TCACTIGTIGGC CTGAGTTCCC CAGCCCCTCT CCTCCTTTTT TTTAGATGAG ATTTAGCACA CICICAGITA TITAAACATG CAACATTICT TGAGTATGTA TGTTGAGGCC ATCTGAGCTC ATAGCTICATT CAGTAACCAG TITICATIGCTIC TGTCATTICAC ACTCACTACT TAATACTIGCC ATGGTGAAAA TGTGGAGGAA AAATGTATCC ATGTGTGTCT GGGAAGCATA TACACTTGTA AAAAAAAGT GATAAAGCAA TCAGAAGACC AAGAGGTTTA CTATTGATGC TTAGGGTCGT CTGACCTTGG CTGGCCAATA GACCTACACG GCCAAATTAA TTTAGGAGAG TAATAATTTT TCAAAAGCCA AITITITITC TGIATITICT GIATGAAACT GCCAATAICA TGAATAGAAA GGGAGAACCA TAAAGGAGAA AGAACGTGAT GTTCTTATT GTTCATGTAA ACCTAAAGAA ACAGITETIGA GGCAGGGGG ATCAGCCGAA CTCTAGGGAC TTGGTGTTGC TTGGAAGGCA ICCATACCIG CATITIGCAT ICTICGIAIG TAAICATAIT GCCAAAGACA AACTAITICA SEQUENCE DESCRIPTION: SEQ ID NO: 134: (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1565 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 134: (X 2 13 8 25 3 35 5 45 တ္ထ 55

TCATTIATIG TAAATAACAC TITTICCCCAG ACCTACCATA AAGTITICIGI GATGIATIGI

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(2) INFORMATION FOR SEQ ID NO: 135:

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Ê SEQUENCE CHARACTERISTICS: TYPE: nucleic acid LENGTH: 2007 base pairs

€ 6 6 6 TOPOLOGY: linear STRANDEDNESS: double

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20 CATTTICTICGC CAGGCTOGAC AAAITTCCTGA GGCACAACTT GGCTTCAGTT CAGATTTCAA TOTALAGEC COOTTATAGE COACTITIONS CAGCAMAGAT COCCONSCAS STEACAGECT 180 120

SEQUENCE DESCRIPTION: SEQ ID NO: 135:

OCTOTOTTOG TOTTOGGACC AGCAGAAGGC AAACGTCCAG CCAACACACA GGACTGTAAG

CTTCCCCAMA GTCAGAGGTG ATTTGATTTG GGGAAGACTG AATATTCACA CCTAAGTCGT AGGACTOTGA GOTACGTGCC CTGTGAAGAC CCCCAGGCTT TGTCATAGGA GGTCGTTCAG

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30 GAGCATATICE TGAGTTTTIAC TICCTTATIGG CITIGCCCTCC AAGTTCTCTC TCTCATACAC ACACACACC TIGCICCAGA ATCACCAGAC ACCICCATGG CTCCAGCTAT GGGAACAGCT

35 GCATTGGGGC TGCCTTTICTG TTTTGGCTTAG GAACTTCTGT OCTICITOTO GCTCCACTOS

TGATCAGCTC GIGIAAAACA CACCGICTIC TIGGCCTCCT GGCAGTICTI GGGCTGCAGG CAGCTCTGGG ACCIOCACACI

TOTOCOAATA GTOCTOTOCO TOGCCAGTTO AATGGGGGAA GCTGCTGGCA CAGGAAGGAG

6 OGCTGAGGCT TAGGAAATTG CTGGAGCCGG CTCCAAGCAG ATAATTCACT

AGCATOTOAA AGTOAAAAGO CATOTOGGOO TGOTGOTTOT OTTTOTOAGG CTOTGGGGAA TCAGAGICAA ACATCATTCT GCCIGTKTTG GGGGCCAGGT GTGTCACACA

3

AGGAATOTOC CTOTOCTOTO ACTIGATIOO AAGTGTOGTT GAATTGTOTG GAGCACTOGG

50 ACTITITITIC TOTTITICCIT GATGGACCAA CAGTGCAAAT GCAATCTCGC CATTTAACTT

ACACCTACGA TOTOCCAGGC ACTOTOTTAG GCGCTTTTAT ATAGATCCTC GTTAGGATGA TCAGGTCGAT TTCCTTTCCT GATCAGACAT CTTTGTGCCC CCTTTAGGAA GGAAAAGAAT

GACTAAGGGA TGAGGACATC TCTTTATAAA AGGCCCCTAA GTAATGGATA AACAGAAAACA

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1140 1080 1020 960 900 840 780 720 660 600 540 480 420 360 300

TOTTCTCAAG GACTTATCCC CTACAATATT CTCCCACTCC ATACTTCTCC TTCTACCCCA CITAGAGOTG AGAAGGTCTG TCTTCAAGAT CCAAGGTAAG ATTOCCTTCA GTCTGATGTT

1260 1200

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(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 136:

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AATTATTTTC TCTGTATGAT TAAAAGT

GTCGCCAAGA CATCTACATT GTAAGAGAAC ACAGTGGAAG ATCCTGTCCT GATTCTCAAA TATANACTOT ATANAAGGIT CIGITITITAA AGGIGGATIT TCATTCCTCT GGGGACAGIG TITITITITA AATITAGGAT AACACATITIT TOITICTAAA GIGATITGIG AITIGIGCIG GARTAGAAAC TGATAGCATT AAAATACICC GITCCICICT CICITCICGC TICCITITII

1980 2007

1920 1860 1800 1740 1680 1620 1560 1500 1440

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CTGTAGCTTT TTAAAAGGAA ACCCAGTCAT CCCACTATGA ATCTGGCATC TTCTTATGCT TIGACAGGIT GOGCIGIGIG TOTOCOCATO TOTOTATACA TITICCAGGCG TOCCIGIGIC TACCTICTOC TIGHTGAGIT GITTIGGCAT ICATATIAAA AGCCAGCAIC ICACIATITA TOGGGACCAA TAGAATATOT GATOTOTGAA TTTTCTTTAA AAAACTTAAG GAGTCTTTOC CCATOTOCTC CCGTGCACTC CTCAGATGGT CAGAGGGGTA ACCCAAGTCC TTAGAGAATT

TTCAGGAGGG GCTGGATCAA ATTTTGAGAG GCTTATGGGA AAGGGAGGGG GAGAAGAAAT TCTAGTOTTT TGGCCATACA TCAACCAAGG GGTTTAATTT ATCCAATGCT TGACGACATG

TGACATTIAT TITATTATIT ATTITAAATG TITACATCTT CITTATGTTG TATCAAGCCT

(A) LENGTH: 1291 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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Ě SEQUENCE DESCRIPTION: SEQ ID NO: 136

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CTTTTNACCC TCCCCCTTCA CACACATACA TATCAGGITG TTTTCTAGTT AAAAACCCAA GIAGCICAGA TICTACTITA AIGICAGIGC AGATITICCAT IGAATCAICC CATTAIGITI 180

TITCICATIT TIATOCIGIT GOGICTIAGT TITTAAATIG ATATAAAGAA CICAGCAATG

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AACTICAATG GICTACIGAA ACAAAAATGG TAACTITICA TIAGIGGATT ATTIAGAGTI GITTIATTIT CTACICATAC TIAGGGITTA GGAAACACIA CCACTAGITA TCATITAATC

TCATACAGIT ATTCCCCATGA AAGGCAGAAT GITTGTTTCA AAATTAATCT AGTTTTCTGT ATAGTAGTIG TITICCAGAAA ACACTICCTC ACAATIGTAC TICCCAATCA AATCATGIGA

> 420 360 300

ATTICACTIT GICTIATITG TAAATAGIGA ACTAAAACIT TIGGCAGAIC AGCAACAITI ACATTIMAAT TIGAGMAGGI GACMACIGGC ICTITICCAG ICTICCTICA IGICAGITTI CTGATAGACC ACTATTOGCA AACAGTATCT GTCAACTACC AAATGTGTAA AATTTTCTGT 480 600

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	361		362
	GCTGAGCCTG THTTTAAGC TAATGTGTAT TCTTACTAAT GTTCCTATCA AGAATGGATT	099	TICHAGHA GHISCACHA CARANTATI GRANTIYAA ACTAAGAANA AAGCCHGCAT
	IGIAATATAT GCIGICTATT ICTAATGITC ACATICATAT TITGAGGITC TATCITATT	720	TITCAMAN AANITOGAAT TOCTOTIGGT GAMADAACAA CCAMANIACT GAATCTGATG
5	TANTAGAGAA CAGACTYCTC AAAAANCTT CAGAAGCAGC TINTIATTGA AATAITCGAAA	780	TROBURCAGO TITICIACAGO AAGAGATGOT ATRATITIACA ATTITICADAT TITAATAACA
	TATTGABATA AACCCGGTGG GTTAGATTAC TCATCTGTCC ACCAAGTGGG ACATTTGCAT	940	CZZZTRACYCE GABABACHCA CHINTARABA WZFRACPAPA BAZRBACZCA WZFRYNYYYY
2	GGACTGGGGG CTTAAAGGAC TTAGAAGAGA CCTGTAAGTA AATCCTGAAA ATGAGCCAAT	01	
2	CCCCACTIGA AIGGITACTG GAGTAAACCC ACCTITACCA CCCCAAITAC AGCACCGGAG	096	
	GCCGATAAAC CAACITGGCT CTGGTTCATT TITICITITICT TCATITIOTGA TGCTCAGAIT	1020	HENCANGESC ATTITITAGNA GEARAGACTC PURGINICISC ATTITICABRICE TRYCELANDED
15	CAAAATGTGT GTTCTACACT GTTACAGGCT TCTCTTTTGT TTGATTAAAG ATTTTAGTCC	1080 15	TITITITICAC AGTIMATOTI COCICCOCAA GITIGOTATI CAAATCAACT GOCTGAANGA
	TACTITISTA TGGACACATT AGAINTTCA GAGACCAAAA TÄGAAGAAIT TGCTGTTHGA	1140	CATTICTAGT AGTCTGATGT ATTTTTCTGA GGAATAGTTT GTGATTCCAA TGCAGGTGTC
70	TATTITICAD ANGICAGCAS ALTIGIGGCA ANICATITIAL TIGCCTITIT ANAMALICAL	1200 20	ITCATTACCA ITACCTCTAC ACTGCAGAAG AAGCAAAACT CCTTTATTAG AATTACTGCA
	TTAAGCAGIT CAGAGAGIAG ACTACTCAGA AAATTATTC AGGTAAITGT CTAAGAGGTC	1260	CANGIGIANG GGGAAAANAG TITCTIGAAAGG CTAGAANGAT ACAAGTIGAG AAAAGTITGGT
	aatattittt aatocatatt gaatcaaata a	1291	
25		. 25	CCTAAACTCC CTTGGGAATC TGGAACAAA GAATATGAAA ATTGCCATTT GAAAACTGAC
	(2) INFORMATION FOR SEQ ID NO: 137:		CAGCTAATET GGACCTCAGA GATAGATCAG CCAGTGGCCC AAAGCCATTT CAAGTACAGA
. 30	(i) SEQUENCE CHARACTERISTICS:	30	AATTATAGAG ACTACAGCTA AATAAATTTG AACATTAAAT ATAATTTTAC CACTTTTTGT
			CITIANAGC ANAITHGIAA ACTCAGAACT GAGCAGAAGT GACITTACIT TCTCAAGTIT
2	(C) STRANDELNESS: double (D) TOPOLOGY: linear	\$6	GATACTEROT TEACTOTICC CITATECCTC ACCCITCCC TICCCTITICC TARGECIATA
તે	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 137;		GTGCACAACT TAGGTTATTT TTGCTTCCGA ATTTGAATGA AAAACTTAAT GCCATGGATT
	GOCACGAGGA CCTACTTITG TAACAGACCA TOGTTGTGTC CAAGGTAAAA CCACAGTGAT	09	TITITICITIT GCAGACACC TOTTIAICAT CITOTITIAAA TOTAAAIGIC CCCTTAIGCT
9	AITITIOGAI GCITIOTOTO CAALCTIGAC ITOTITITIGO AGEATCATIA ITCACACTIC	120 40	TITICAARTAA ATTICCITIT GIAABAAAA AAAAAAAA AAAAAA
	AAATIGIGAA TCITITIAAAC AICTIGARAA TITIGITIGITIG AGAGCIGITC AITICIAAAAT	180	
45	GTAATGAAAT TCAGTCTAGT TCTGCTGATA AAGATCATCA GTTTTGAAAG GTTACTGATT	240 45	(2) INPORMATION FOR SEO ID NO: 138;
	ITCCTCTTCC CTCTTAGITT TITACCCAAT AIAIGGAGAA GAGTAATGGT CAATCTTAAC	300	(i) SEQUENCE CHARACTERISTICS:
	ATTITICITIT AATTICITIAA TAAAGCIGCT GOGCAGIGGI GCAGCATICC TACCTAGIGI	360	(A) LENGTH: 1935 base pairs (B) TYPE: michele and
20	CATRADAGCA ADATRCTTAC ATROCTITICT TADARTATRG GAATGACATT ACATTITTAG	420 50	(C) STRANDENESS: Gouble (C) TOPOLOGY: linear
	GAGADAGTAA GTTGCTTTGC ACCGCCTACT TAATTCCTTT CCATATATTG TGATACADAC	480	(x1) SEQUENCE DESCRIPTION: SEQ 1D NO: 138:
55	TITIGAATAT GGAATCTIAC TATTIGAATA GAAATOTGTA TOTATAATAT ACATACATAC	55	POTGRACTAA TECTRACAGA POCCOCTIGAG GGATTOCTTGA TOGGCTGAGC AGCTGGCTTGG
	ATAAGCATAT ATOTICTICT CHOTOTICTAT ATATATATA ATGCATGCTG TGAAACTTGA	009	ACCINGIACT GACTGACATT CATTGTGATG AGGGCACTT TCTGGTACAG GALTTCTAAGC
	CTACACAACA TAAATCACTT TITAAATTCC AGGAACGGGT AGTCTGACAC GGTGATTATC		TCHNISTETT ATARACATT TCATCISTAC TTSCACCTCA CTTTACACAA CACGAAACTA
9	CITITICAGGC TGAATCCGIT AITAACTIGI TATITAGGIT ITACICCCAG TAGCAAGGGA	720 60	

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TOCAMAGITA GCTOGATOGC TOMAGGTOMC TIMOGTAMOI TOGCAMGICC MISCITICOCA

300

TAATOGATAA TOTTATTTAT OTAAACTAAA GOTTOOTOTT TATACACACT COTOTTATTO CACTAATIGG CTAIGICICT GGACAAGITT TITITITITI TITITITIAA ACCCTTICIG GCAGAGCAGC AGTTAAACCC GTGGATTTTG TAGTTAGGAA CCTGGGKTCA AACCCTCTTC AAACACTATC TICCCATCIG TITCICAATG CCTGCTACTT CTIGIAGATA TITCATTICA CAATAGCATG AGGTGAACAG GACGTAGTTN AGGCCTTCCT GTAAACAGAA AATCATATCA OTTICIOCCT CTITITIGGAA AAGAAAACAA AGIOCAATIG TITITITACIG GAAAGTIACC STIGCTIAGET TIGGAAAGIC TAGAATGGGT CCCTGGTGCY TITTIACTIT GAAGAAATCA CTCAGCTCCT CAGGTCAGCA AGTCTACTTC TCTGCCTATT TIGTATACTC TCTTTAATAT TIGCTATICG CAGCTACATA CAACGIGGCC AACCCCAGIA GGCIGATCCT ATATATGATC TOGGATAAGA TAAATGACCA CAGTACCTTA ATTICTAGGT GOGIGCCIGT GATGGTICAT TAAAGOOTET GECAGATAGG AAGATGETAG TTATGGATTT ACAAGGTTET TAAGOETGTA GAGGGGGGTA GCATGCACCC AGCAGGGGAC TGAACTGGGA AACTCAAGGT TCTTTTTACT TOTAGOTANG GACATTTTCT YTTTTTCNGC AGCTOTOTNG GTCCAGAGCC TCTGGGAGAG AAAACTAGCC AGAAGTCTCT TTTTCAAATT ACTTACAGGT TATTCAATAT AAAATTTTTG AGAGTETAAA ACETACAGTG AATCACAATG CATTTACCCC CACTGACTTG GACATAAGTG AACTTICACI TICIATGICI ACCICAAAGA ATIGITGIGA GGCTIGAGAT AATGCATITG GICGCACAAT AGGCCGICIG CAAGCIGGGI TAGAGAGAAG CCAGIAGIGG CICAGCCIGA TGAAACTTAG AGTTATAATT CATGTATTAG GGTTCTCCAG AGGGACAGAA TTAGTAGGAT AATTAGGCAA CCTAAAATAT TGATGCTGGT GTTGGTGTGA CATAATGCTA TGGCCAGAAC AGTGCTGGTG CTGACTCTCA ATAGCCCCAC CCAAGCTGGC TATAGGTTTA CAGATACATT GIGGGGIAGI GAGCIGCCII ICIGIGAICG GITICCCIAG GGAIGIIGCI GIICCCCICC GITCAAAAAC CICAAAACIG GGGAAGCIGA CAGIGCAGCC AGCCIICAGI CIGIGGCCA ATATOTATAT ATGAAAGGGA GGTTATTAGG GAGAACTGGC TCCCACAGTT AGAAGGCGAA CTTTCCATTA CATGAGCIGT CTCAAAGCCC TCCAATWAAT TCTCAGTGTA AGYTTCAAA AACCTOGAGT CTGATOTCCA AGAGCAGGAA GAGTOGAAGA AAGCCAGAAG ACTCAGCAAA ACCICAAGAG CCCCTGGCAA CCAACCCACT GGTGCAAGTC CTAGATTCCA AAGGCTGAAG TECCTGAAGT TOCCCTOGTE TETGCACCTT CTAAACCTAG AGTOTICTACC ACCAYAGTOG CCATACCAAA GAGGCTACCG TTCTTAAGAC ATTCCTTCCT 1200 1440 1320 1260 1140 1080 1020 1920 1860 1800 1740 1680 1620 1560 1500 1380 960 900 840 780 720 660 8 540 480 420 360 1935 25 20 5 3 8 ႘ છ 5 S 50 55 8 TOGATIOTOG TITTCCCCAGG CAGAGGGCCC CICTYTTCCC AGCACTICCC TGCCTCCCCC TODOTAGOTO GATTTOAGCO GAAAGACTCC CAAGAATOTGC CAAGAATTTC CCROTCCCAG GGAAGCTTAT TTTCCCGTGG CCAGGATGCA TTTCTCTGAG TGGAAACAGG TTCTTGCATG GAGGACAGAC ACAGATTTCC TOCTGGGGGA GGGAGGAGTC CACOCATCCT GATOCTGCCT NOCCCCCTTG GCACAAGTCA GATGAAGCAC GTTCTGCCGG GGAGGCCCTC AMCTTCCAGA (2) INFORMATION FOR SEQ ID NO: 139: GUAGUAGUTG AGGAGGGTGA ACAAACCCCCG AGGGAGGCCC GGCCCTTGCT CCCGAGTTGG CAGGASTAAT TIGAAATOTG TGAGGTGACT CCCCGGAGGC CTTGGGCTTG GGCATTTGGG TCAATGOTGO AGGGAGGCAG TGGGGGAGAAA GOTCTCACCG GACAGOTTGG GAGAATGAGG CICCICAKOT GACCACCCIC CICIGACCSA OGCCCCCICC ITGICIGAAA AAAGGAGCCI TECCETTEAT CTECOTECCE CTETTTGAAG CTOTECCEAT CTEAGTOTEA GACCAGCETT GOCTECCOCC TOTOCCCCTT CICCTCACCA TOTCTCCCCC ACCCTGCCTC GCAGGOCAGG GGAAACTAAG GGCAAGCAGG ATACAGGOCG AGGGATGTGG CAGGTGAGGG ATTOTOGTCC GAGGACAGCC TITTCCCCAG OCCTCARAGC ATTOCTCATC OGTOCCAAAC GOGGAGGGG TOTOGCAACG TOCOCCCCGC AGAGGCCACG CATOTITIGAC CAAAGCCCTC ACOCCTCAGO CCAGCACCCA GITCCTCCTC ACATGOCAGG TGAGCACAGA CITCTAGITG TACCCCAACC CTTTGCGGCT CTGTACACAT TTTTAAACCT GGCAAAAGAT GAAGAGAATA CTGTHAGACAG ATGCCCTCAG AATTGGGGCA TGGGAAGGGGG GSTGGGGGAC CCCATGATTC GGICTAGCTC CTCTCCAMAC AGCCATGCCC TCCAMATGCT AGAGACCTGG GCCCTGAACC CTUACACCCA ANICCAGAAT CCCTGGTCTT GAGTCCCCAG AACTTTGCCT CTTGACTGTC AAAAGAATGA TOTCTOGAAG GGCTTAAGGG ACACAGTGGA COAGGGGAGA GTCCTCATCT TCHGCGGTGC TGGGGHACAG ATGGAGGGGG CAGTGGGGAC AGGGCTTGGG CAGACACCAG COTTOTOTO CTACCTOCAT COATGGAAAA TTAGTTATTT TOTGATOOTT TOOCCTGOOT AGCCACGGAC TCCAATGCCC AGCTCCTCTC CCCAAAACAA TCCCGACAAT CCCTTATCCC (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139: TOTOGOGIST TAGTOCCANA CTIGANTAGG GGCTGGGGTG CTGTCTTCCA (A) LENGTH: 1446 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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960 900 720 660 600 540 480 420 360 300 240 180 120

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1320 1260 1200 1140 1080 1020

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WO 98/39448

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10 (1) INTORNAL TO SEQ ID NO: 140: (1) SEQUENCE CHARGETERISTIES: (1) SEQUENCE CHARGETERISTIES: (1) THE SEQUENCE CHARGETERISTIES: (1) THE SEQUENCE DESCRIPTION: SEQ ID NO: 140: (1) THE SEQUENCE DESCRIPTION: SEQ ID NO: 140: (1) THE SEQUENCE DESCRIPTION: SEQ ID NO: 140: (2) THACCHARA ACTIVITY THE ARTHOUGH ACTIVITY TO CHARGE ACADICATA THE ACADICATA ACTIVITY THE ACADICATA CHARGE ACADICATA THE ACADICATA ACTIVITY CHARGE ACADICATA THE ACADICATA ACADI	WO 98	WO 98/39448 . PC	PCT/US98/04493	WO 98/39448	PCT/US98/04493	5
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(i) SEQUENCE CHARCTERISTICS: (ii) LENGTH: 1109 base pairs (ii) SEQUENCE CHARCTERISTICS: (ii) TUBENTH: 1109 base pairs (iii) SEQUENCE CHARCTERISTICS: (iii) TOPOLOGY: Linear (ivi) SEQUENCE DESCRIPTION: SEQ ID NO: 140: THITTITITIT TITICATHER ANTOTICTT CTCCAFTECA CHARTHAGCT AGGRANCHC THACCCAMA ACTITICTA AGGRETICC TITICAGGGCA CHARTHAGCT AGGRANCHC THACCCAMA ACTITICTA AGGRETICC TITICAGGGCA CHARTHAGCT AGGRANCHC THACCCAMA ACTITICTA AGGRETICC TITICAGGGCA CHARTHAGT ATTENNOCY ANTOTICAGA AGGRETICAGA AGGRETICT TIMOTICGA CHARTHAGT CANCENTRY THACTCCAGT ANTOTICAGA AGGRETICAGA AGGRETICT TIMOTICGAGA AGGRETICAT THACTCCAGT ANTOTICAGA AGGRETICAGA AGGRETICT TIMOTICGAGA ACTOCAGT ANTOTICAGA AGGRETICAGA AGGRETICT TIMOTICGAGA ACTOCAGT CHARTHAGA AGGRETICAGA AGGRETICT TIMOTICGAGA AGGRETICAT TIMOCAGAA AGGRETICAGA CAGGRETICAT TIMOTICCTT TOTICAGAGA AGGRETICAT TOTICGAGAACT TICCTICAGA AGGRETICAGA AGGRETICAT TOTICGAGACATA CAGGRETICAT TIMOTICCTTA TIMOTICTATA AGGRETICAT TOTICCCTTG TCACTITICTT CAGGRETICATA AGGRETICAT TITICCCCTTG TCACTITICTT CAGGRETICATA AGGRETICAT TITICCCCTTG TCACTITICTT TAMITITICT TAMITITITT TITICCCTTA TTCTTCCTTTTTAGA CAGGRETICTA AGGRATICA AGGRACATAC CAGGRACATA TTCTTCTTTTTAGA CAGGRATICA ATTCCTTATA AGGRATICA AGGRACATAC CAGGRACATA ATTCTTATA AGGRATICT AGGRATICA AGGRACATA TTCTTCTTTTTAGA AGGRATICAT ANTOTICCTA ATTCAGAGA AGGRACATA ATTCTTATA ATTACAGAATTC CAGGRATICA TCTTTTTAGA AGGRATICAT ANTOTICCTA ATTCAGGAAT TCTTTTTAGA AGGRATICAT ANTOTICCTA ATTCAGAGA AGGRAGACA ANTOTICTA TAMICTAGA ATTCAGAGAA AGGRAGACA ANTOTICTA TAMICTAGA ATTCAGAGAA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGAA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGAA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGAA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGAA AGGRAGACA ANTOTICTATA ANTOTICCTA ATTCAGAGAA AGGRAGACA (1) SEGUENCE CHAGACTERITATA ATTACAGAA (1) SEGUENCE CHAGACTERISTICS:		A COMPANY AND STATE OF THE STAT		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 141:	 	
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AGTICOGOAGO ANGICTITAT TCTAATOTOA GGGIAGGGAA AATGICGATA ACATTACTOG GGTGARGGAG GCATTGITCT TIAGTICGAG TTCTCATTIT TATTCTCCAG TACTGACTTG TGGGGAAAGC ATACTITITC ACTGCCAGGT ACTGAATGCA GAGGTTATAA TGTGGGAAAG CATACTITITC ACTGCCAGGT ACTGAATGCA GAGGTTATAA TGTGGGAAAG GGGTTAAAG CCTTAAGTGA ACAAATCTAG CTAACAGTGA ATGAACTAGT GGTTTTGGAAA GGGGTTAAAG CCTTAAGTGA AAAATCTAG CTAACAGTGA ATGAACTAGT TAGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG ACCAATGTTT TGTGAAAGTG GTTTATTCCA CTGGAGAAAG GGGATTGAAA ATCAGTTAGA ACCAATGTAT TTCTTGCCCC ACGGAACACT ATTCCTATAA GATAGCAGTGA AGAAGCTGCT GTGAGGAGCT CAGCTCCCAAA CACAGGATCA ATATCTCTATAA GATAGCAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTAT CAGAATCCC TAAGCATTTT AATAAAAAGT TAAAACAAA AATAAAAAGG GACACTCGAG, GGGGGGCCCG AAACCCAAT (1) INFORMATION FOR SEQ ID NO: 141: (1) SEGUBACE CHARACTERISTICS:		ttataataa aagatcaaaa gatatatctc ctatgaacag attggagata ggagatgada	300 25	AAAGAATTA TCTAAAA	497	
GOTGARGAGA GCATTGTTCT TTAGTTGGAG TTCTCATTTT TATTCTCCAG TACTGACTTO TGGGGAAAGC ATACTTTTC ACTGCAGGT ACTGAATGC GAGGTCAGT GAGGTTGTT GGTTTGGAAA GCAGTTAAAG CCTTAAGTGA ACAAATTCAG CTAACAGTGA ATGAACTAGG TAATATAACT TGCATATTTT TAATTTCCTT TGGTTAAAG CTCCCCATAC TTCTCTGTTC GGAGACATGA GAAGTTAGAT TACTTCAGTG ATCTGTGGG TTATCATTTTT TTCCCCTAT TTGTCCCTTG TCACTTTGTT GCAAGCTAGA ATCTGTGGG TTATCACATAG GGCAGCTCTT TTGTCCCCTG TCACTTTGTT GCAAGCTAGA ATCTGTGGG TTATCACATAGA ACCAATGTAT TTGTTGCCCC ACGGAACA TACTCTATAA GATAGCTGAA ATCAGTTAGA ACCAATGTAT TTCTTGCCCC ACGGAACA TATCCTATAA GATAGCTGAA ATCAGTTAGA ACCAATGTAT TTCTTGCCCC ACGGAATCA CTCCAGAAAT AATAAACAAA ACTAAAACAAA ACTAAAACAAA AATTAAAAG AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAAT TAAAACAAA AATTAAAAG AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAAT TAAAACAAAA AATTAAAAG AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAAT TAAAACAAAA AATTAAAAG (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:	¥	CITOGGAGO AIGICITIAT ICIAATOIGA GGCIAGGGAA AAIGIGGAIA ACAITACIGG	360			
TGGGGAAGC ATACTITITC ACTGCAGGT ACTGANTGC GAGGCTCAGT GAAGTATTA TGTGGGAAGT GCATGCTTT COTTTATTAG CAAACTGG TGGATTAGG CAAAGTTGTT GGTTTGGAAA GGGGTTAAAG CCTTAAGTGA ACAAATCTAG CTAACAGTGA ATGAACTAGG TAATATATAACT TGCATATTTT TAATTTCCTT TGGTTAAAG TCCCCCATAC TTTCCTGTTC GGAGACATGA GAAGTATTT TAATTTCCTT TGGTTTAAG TCCCCCATAC TTTCCCCTAT TTGTCCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGAAA ACAAGTTGTT TGTGAAAGTG GTTTATTCCA CTGGAGAAG GGGATTGAAA ACAGTTGTT TTTCCCCC ACGGAATCATAA GATGCATGAAA ACAAGCTGCTT TCTCTTCCCCC ACGGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTCTTTTTTT CAGAGTCCAT ATATTCCCTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTTAT CAGAGTCCAT ATATTCCTA ATAAAAAGT TAAAACAAAA AATTAAAAGG GACACTCGAG, GGGGGCCCG AAACCCAAT (1) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:		STGARGAG GCATTSTECT TIADTICCAG TECTCATTIT TAITCTCCAG TACTGACTTG	420	(2) INPORMATION FOR SEQ ID NO: 142:		
GOTTTGGAAGT GCATGCATTT COTTATTAG CAACATAG CTAACAGTA ATGAACTAG GOTTTGGAAA GGGGTTAAAG CCTTAAGTGA ACAATCTAG CTAACAGTGA ATGAACTAG TAATATAACT TGCATATTTT TAATTTCCTT TGGTTAAAGG TCCCCCATAC TTCTCTGTTC GGAGACATGA GAAGTTAGAT TACTTCAGTG TTAGTTTTCT TAATTTTTT TTCCCCTAT TTGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCAGCTCTT TTGTCAAAGTG GTTTATTCCA CTGGAGAAG GGGATTGAA ACAGTTAGA ACCAATGTAT TTGTTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT CAGCTCCAAA CACAGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTAT CAGAATCAC TAAGCATTTT AATAAAAAA AGTAAAACAA AATTAAAAGG GACACTCGAA, GGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUENCE CHARACTERISTICS:		330GAAAGC ATACTTTTTC ACTGCCAGGT ACTGAATGCA GAGGCTCAGT GAAGTATATA		(1) SEQUENCE CHARACTERISTICS:		
GOTTIGGAAA GGGGTTAAAG CCTTAAGTGA ACAAATCTAG CTAACAGTGA ATGAACTAGG TAATATAACT TGCATATTTT TAATTTCCTT TGGTTAAGG TCCCCCATAC TTCTCTGTTC GGAGACATGA GAAGTATGTT TACTTCAGTG TTAGTTTTCT TAATTTTTTT TTTCCCCTAT TTGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCAGCTCTT TGTGAAAGTG GTTTATTCCA CTGGAGAAG GGGATTGAAA ATCAGTTAGA ACCAATGTAT TTCTTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT CAGCTCCAAA CACGGAACA ATCCTATATA GATAGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTTAT CAGAATCAC TAGGAATTCC CATGAATTAT GACTTCTCAT AGAAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAA AATTAAAAGG GACACTCGAG, GGGGGCCCG AAACCCAAT (1) INFORMATION FOR SEQ ID NO: 141: (1) SEQUENCE CHARACTERISTICS:	¥	GIGGGAAGT GCAIGCAITT COTTIAITAG CAAACAIAGC IGGAITAAGA CAAAGTIGIT	540			
TAATATAACT TOCATATTTT TAATTTCCTT TGGTTAAAGG TCCCCCATAC TTCTCTGTTC GGAGACATCA GAAGTAGAT TACTTCAGTG TTAGTTTTCT TAATTTTTTT TTTCCCCTAT TTGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCACTCTT TGTGAAAGTG GTTTATTCCA CTGGAGAAAG GGGATTGAAA ATCATTAGA ACCATGTAT TTCTTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT CAGCTCCAAA CACAGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTTGTTTTAT CAGAGTGCAT ATATCTCATA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTAT CAGAGTGCAT ATATCTCATAT ATAAAAAAGT TAAAACAAAA AATTAAAAGG GACACTCGAA, GGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUENCE CHARACTERISTICS:		CITIGGAAA GOOGITAAAG CCTTAAGTGA ACAAATCTAG CTAACAGTGA ATGAACTAGG	600 35	(c) Siranbinass; gauste (D) TOPOLOGY: linear		
GGAGACATCA GAAGTATGAT TACTTCAGTG TTAGTTTTCT TAATTTTTT TTTCCCCTAT TTGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCAGCTCTT TGTGAAAGTG GTTATTCCA CTGGAGAAAG GGGATTGAAA ATCAGTTAGA ACCAATGTAT TTCTTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT CAGCTCCCAAA CACAGATCA ATACTCTATA TAGAATTCC CATCAATTAT GACTTCTCAT TCTGTTTTAT CAGAGTCCAT ATATGTCCTA CTTCAGGAAA AGTAAAACAA AATTAAAAGG AAGAAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAA AATTAAAAGG GACACTCGAG, GGGGGGCCCG AAACCCAAT (1) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:	2	ANTARACT TOCATATITY TAATTICCIT TGGTTAAAGG TCCCCCATAC FICICIGITC	099	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:		
TIGTCCCTTG TCACTTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCAGCTCTT TGTGAAAGTG GTTTATTCCA CTGGAGAAAG GGGATTGAAA AGAAGCTGCT GCAGGAGGT TTCTTTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT CAGCTCCAAA CACAGGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTAT CAGAGTCCA ATATGTCCTA CTCAGGAAA AGTAAAAACA TCATTTAGA AAGAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAA AATTAAAAAGG GACACTCGAG, GGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:		GRONCHIGA GANGINIGAT INCTICAGIG TINGITITICE INAITITITIT ITTICCCCINI	720	ATCHGGCAGA GOLARGCTGC CTGCCAACCC CCTCCCTCAA GGAATGGCCT TGCCCAGGAA	09	
TICTICAMAGIG GITTATICCA CIGGAGAAG GGGATIGAAA ATCAGITAGA ACCANIGIAT TICTICCCCC ACGGACACT ATTCCTATAA GATAGCTGAA AGAGCTGCT GTGAGGAGCT CACCTCCAAA CACAGGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGITTTAT CAGAGTCA ATATGTCCTA TAGGAATTCC CATGAATTAT GACTTCTCAT AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAA AATTAAAAGG GACACTCGAG, GGGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:		TOTCCCTTG TCACTTGTT GCAAGCTAGA AATCTGTGGG TTATACATAG GGCAGCTCTT		TGCCACCAC ACATACCTC TTCTTTTTT CTAGTCAAAC TCTTGTTTAT TCCTTGGCTT	120	
TTCTTGCCCC AGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT CAGCTCCAAA CACAGGATCA GCACCTTGTA TAGGAATTCC CATGAATATT GACTTCTCAT TCTGTTTTAT CAGAGTCAT ATATGTCCTA CITCAGGAAA AGTAAAACAAA AATTAAAAGG AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAAA AATTAAAAGG GACACTCGAA, GAGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:	¥	GTGAAAGTG GTTTATTCCA CTGGAGAAAG GGGATTGAAA ATCAGTTAGA ACCAATGTAT	840	GCCTCCCTC TITCCTCCCC TCTCAACCTT TTACTTCTGG TTTCTATTTC ATGGGATTTG	180	
CAGCTCCAAA CACAGGATCA GCACCTIGTA TAGGAATTCC CATGAATTAT GACTTCTCAT TCTGTTTTAT CAGAGTCCAT ATATGCCTA CTTCAGGAAA AGTAAAACAG TCATTTAGGA AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAAA AATTAAAAGG GACACTCGAG, GGGGGCCCG AAACCCAAT (2) INPORMATION FOR SEQ ID NO: 141: (1) SEQUEAVE CHARACTERISTICS:		TETTGCCCC ACGGAACACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT	900 45	OCOTICAMET TAAMETTACA ACAGTOCCOC CAACACCAAG TETTOCAGGA AAAAAATACA	240	
TCTGTTTAT CAGAGTOCAT ATAIGTCCTA CTTCAGGAAA AGTAAACAG TCATTTAGA AAGAAAGTCA ATCTGTATC TAAGCATTTT AATAAAAGT TAAACAAAA AATTAAAAGG GACACTCGAG, GGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:	ฮ	ASCICCAAA CACAGGAICA GCACCTIGIA TAGGAAITICC CAIGAATIAT GACTICICAT	096	aagaaattta acaaalaaaa aaaaaaaa		
AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAGT TAAAACAAAA AATTAAAAGG GACACTCGAG, GGGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEAUC CHARACTERISTICS:		CTOTITIAT CADAGROCAT ATAIGTCCTA CTTCAGGAAA AGTAAAACAG TCATTTACGA	1020			
GACACTCGAG, GGGGGCCCG AAACCCAAT (2) INFORMATION FOR SEQ ID NO: 141: (1) SEQUEACE CHARACTERISTICS:		AGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAGT TAAAACAAAA AATTAAAAGG		(1) THEOREMAN TO PROPERTY (1)		
(2) INF	ਰ	ACACTICGAG, GGGGGGCCG AAACCCAAT	1109	(2) INFORMATION FOR SEQ ID NO: 143;		
(2) INFC			\$\$	£1		
	J	2) INFORMATION FOR SEQ ID NO: 141:		(C) STRANDENESS: double (D) TOPOLOGY: linear		
	09	(1) SEQUENCE CHARACTERISTICS:	09	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 143:		

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TATOGROCIG GGGITTAACA CTAMAAACIA GAAATAAACA TCICAAACAG TAAAAAAAA GACAATCAAA AACGACAACA AGCTTCTTCC CAGGGTGAGG GGAAACACTT AAGGAATAAA GCACTOTTAG GOTTGOTTAG TOTACAAGGG ACAGTTGCAT TIGITGAGAC TITAATGGAG GCTGCTGTGG GGAAGAAGAA GAAAGATGTG AFCCTGGCTG ACTTACTGCC TYCCTYGGCT CTIGCCCTCCA TGATCAAGAG GAGGCAAGGC CACATTGTCG CCATCAGCAG CATCCAGGGC AAAAGGCAAA CTATAGGATA ACACAGAGCC CTTTTTGAAA ATAAATTGGC ATTGGAGTGT GTTTATETTE GAACTETGGE TECTGGGETE TTETTEAGEE TEATGEETEE AGGGEGAGAA STTATOGACA CCACCACAGC CCAGGGCCGA AGCCCTGTGG AGGTGGCCCA GGATGTTCTT GCCTACATCC ACACCAACCT CTCTGTAAAT GCCATCACCG CGGATGGATC TAGGTATGGA ANGATGAGCA TICCITITICG ATCAGCATAT GCAGCCTCCA AGCACGCAAC CCAGGCTTIC GTOGACAAGA GOGTCATOGA GACAAACTAC TTTOGCCCAG TTGCTCTAAC GAAAGCACTC GCTTCTCATG CCACCAAGGT GCAGACACAC AAGCCTTACT TOGTGACCTT CGACCTCACA AMACTEGTEC TCTOTEGCCG GAATEGTEGG GCCCTAGAAG AGCTCATCAG AGAACTCACC ATTTOTOTOA CAAGTOOGAA AGACTGAAGA AACACATOTO GTOCAGATOT GOTOOCAGAG ANGAGCGGAA ATCCANGAAC TCCTAGTACT CTGACCAGGC AGGGCCAGGG CAGAGAAGCA TITIGACTOTO TOCOTOCOGA GATOGAACAG TATGAAATTG AGGTGAOOGT CATCAGOOOC ATACTITOTICA ACAATOCIOG GAICAGCIAC COTOGTACCA ICATOGAÇAC CACAGIGGAT TTCCGGCTGC TGCAGTGGGT GCGCGGAAG GCCTACCTGC GGAATGCTGT GGTGGTGATC TICATCACCI CCACAGCCAT CCIGCCCCIG CIGIICGGCT GCCIGGGCGT CITCGGCCIC ITGATIGACT ATGGICICIC COGCTACCAG GAAGAGICIG CCGAAGIGAA GGCCAIGGAC CCATAGITISC AGCAGCAGCT GAGATCCTGC AGTGCTTTGG CTATGTCGAC CCTCAGGGCT GGGCAAAGAA TGTGCAAAAG TCTTCTATGC TGCGGGTGCT 1269 1260 1200 1140 1080 1020 960 840 900 780 720 660 600 540 480 420 360 300 180 8 120 30 25 20 6 35 15 5 ß 8 25 S CCTTTTGTCT CCTAAAGATA GGGATCTACT TTTGAAGGGA ATTGTTCCTC CCAAATAAAT GTCCCAAAGA AGAGCATGGG TGGCAGATGG TAGGGAATTG AACTGGCCTG TGCAATGGGC TITACCCICT AGCIGITITA CITAGAANGT AACANAIGCT GCCTACCCAC CICAAAANGT TOCAGGITCC GTATTCATAC ATTTCAGCTG CAAGTCAGCA ATTTCCCAGG TACCATGTAA GCTATAAAAC GCTTGACCTC CAAGTAGAGC TGATACAGAG ATCTGTGAAT ATTGTGATAG GICCIGAACC IGGICCIGIG GGCCATIGAA AAGITAGAIC IGIGAICICI GGGGIITITIG CAGACCAGCT TITIGCACAAT GAAGCGCAAG GGAACAAGTG GITTIGCCTGG TGTCCTACCT CCCCCTTGCC TOGTTCCAGC TGTCAGAGGG ATACCATCCT AGGGTCTGGG AATCCAAGGC CAAGITATTY TOATGITACO TGGAGAGIGT COAGAGGOIG CICTGAGGOT GAGGIGIGIT TCACATTIOG CCAAACCIGA CACIGICITG CIGCATICIC CITIGGCAAA CAICAGGGIC TIGOTITIACO TIGGICOTITI CITITIGIGCO AGIATICAAG IGGIATAGOI CIGAGOAGG GCAGAATGGG GTTGGGCCTG TGGCCCCCAA ACTAGGGGGT GTGGGTTCAT CACAGTGTTG TCACCTTTAT TCTTGAAACT GAGGTTTACC TGGATCTGGC TACTGAGGCT AGAGCCCACA AGAACATOOT CTTCTTGCCA CGGGGTGTTG TTGTCTCTGG TGGTGCTGCA TGTCTGTGGC ATGGAGCACA AGGGGTCACA GCATGCCTCC TGCCTTACCG TGGCAGTACG GAGACAGTCC CTOTACTOCA AGAGGGCCCT GGGCCTCTGC TTTCCATATT CACGTTTGGC CAGAGTTGTA AACAGTGGGG ACTUATION TOOCTITIGIT CAATGCTTCC ACTCTAGGGC AGGCAGAGCA GTCTATACTC CACGAGACTC CTTGGTTTGT GGTCCGAGAT CCTGTACTAA GGAGGGTCTG GCCAGAGGAA GOCCIOTOTA GAGOCCCCIC CIGIOCCCIC AGIGGCIGIC GITTIGITAAC AICAICAGA GIGAATOCTG AGAATICAGG ATAGECETIC CTAGGGEACT GGACTITETG GEATGGGGGE TGTGTTTGEA GIGGECETGG ATOCTGGAGE TGGAGGGTTT TETGTGETEA GACTGTAGEE TGTAGETETT AGAGACTOTO GTATTITICAT GAATTTACCA TATATCTITG TITTITCTICA ACGAAAAAGT TAATTGAGGC AGATGOGAAA GOTCAGOCAG AATTTTTCTG CCCTACAAAG GOTGGAAGAG AAAGGACACA AAAGACAGAG GATAGCTGTG GGGCATTGTC TGGGCTGCAG GGCTGCCAGG TTCTGTACTT GTGTCCAGCT CCIGIGATAC IGCCATGGCA CAGGATCIGA GITGCAGCTC IGCACCCTAA TATTOGCACA GGGANTAANT ATCCTGGCGT CTGGAGCCTT CACCTCTCCG AGCTAACTOT GAATGAGGAG AGTACCTGCT GCAGGACCTG GAGGTCAGGT GOGTOGATTG GGAGAACAGA TTAGGAGTAT AGCAAATGAA CCCAGAATGG CTITITICCIT CCICAGGITT reference referrence ACTICATOGGA TICATGAGGTIC CCAGCAAGGG CATGGCTGGT AAATTCTTTG TCCCAAGCCT

840

780

720 660 600 540 480 360 300 240 180

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960 900

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PCT/US98/04493

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WO 98/39448

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SEQUENCE DESCRIPTION: SEQ ID NO: 144:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1944 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(2) INFORMATION FOR SEQ ID NO: 144:

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1380 1320 1260 1200 1140 0801 1020

1500 1440

1680 1620 1560 8

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369		370
AATGICAICT GCICAAAGIT GAOIGOIJIA TICACAAJAA ACIGIAAGIT TCIGAITATA	1920	(B) TYPE: nucleic acid
ANANAANA MAANAANA ANG	1944	(D) TOPOLOGY: linear
	8	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:
		GSCACGAGGA GGGCCACGGC AGCCATCGCG CTTTGCAGTT CGGTCTCCTG GTGTACGGCC
(2) INFORMATION FOR SEQ ID NO: 145:		ANCICCANOT AGGGGATTISC GITCCCTCCA GTCGCAGACC CTATCAGATT TGGATATGTC
(i) SEQUENCE CHARACTERISTICS: (A) LENTH: 101 base pairs	01	CTICATAITT GAITIGGATTT ACAGTOGITT CAGCAGTOTG CTACAGTITT TAGGATTAIA
(B) TYPE: nucleic acid		TARGAAAACT GOTAAACTOG TATTTICO ATTIGGATAAT GCAGGAAAAA CAACATTICTT
	15	ACACANGCTA ADACANGACA CACTINGGACA ACANGINGCOA ACATITACANG CYACTIONTA
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:	1	AGAACTGACC ATTGCTGGCA TGAGGTTTAC AACTTTTGAT CTGGGTGGAC ATGTTCAAGC
TCGACCCACG COTCCGGGGT GCGCAACGGG GAGTTCCGGC TGGAGACCCG TGCTCTGGGC	09	TCGAAGACTG TGGAAAACT ACCTTCCTGC TATCAATGGC ALTGTATTC TCGTGGATTG
COGCOCCTIC ACCATGOCCT COGCAGAGCT GGACTACACC ATCGAGATCC COGATCAGCC	120 20	TOCHORCEC GAAAGOCTOT TAGAOTCAAA AGAAGAACTT GATTCACTAA TGACAGAAGA
CTGCTGGAGC CAGAAGAACA GCCCCAGCCC AGGTGGGAAG GAGGCAGAAA CTCGGCAGCC	180	AACCATTGCT AATGTGCCTA TACTGATTCT TGGGAATAAG ATGGACAGAC CTGAAGCCAT
TOTOGTGATT CTYTTOGGCT GGGGTGGCTG CAAGGACAAG AACCTTGCCA AGTACAGTGC	240 25	CASTGAAGAG AGSTTOCGAG AGANSTITTOS TITANIANGST CAGACAACAG GAAAGGGGAA
CATCTACCAC AAAAGAGACT GCATCGTAAT CCGATACACA GCCCCGTGGC ACATGGTCTT	300	TATATCTCTS AAAGAACTSA ATOCCCSACC CTTAGAAGTT TICATGTGTA GYSTGCTCAA
CITCICCGAG TCACTGGGTA TCCCTTCACT TCGTGTTTTG GCCCAGAAGC TGCTCGAGCT	360	AAGACAAGOT TACGGAGAAG GCTTCCGCTG GATGGCAAAG TACATTGATT AACACAAAACT
GCTCTTTGAT TATGAGATTG AGAAGGAGCC CCTGCTCTTC CATGTCTTCA GCAAGGGTGG	420 30	CACATTOSTT CCAGGTCTCA ACSTTCAGGC TTACTCAGAG ATTTGATTGC TCAACATGCA
COTCATOCTG TACCOCTACG TACTGGAACT CCTGCAGACC CGTCGCTTCT GCCGCCTGCG	480	TAACTIGAAT TCAATAGACT TITGCTGGTT ATAAAACAGA TGTTTTTAG ATTATTAATA
TOTOGROGIC ACCARCTITO ACAGOGOTOC TOGROADAC AACCTOGRAA GGOOTOTOCO	540	TTAANICAAC TTAATTITGAA TGAGAATTGA AAACTGATTC AAGTAAGTTT GAGTAATGA
GOCCETOGCA GCCATCETIGG AGGCCGGGC COCCATGCTG CGCCTGTTGC TACTGGTGGC	009	AIGITAGOTT ICIDARITICA TRABAGIROT INSCITITIRE ACITITADART CINGACATOR
CTTTGCCCTG GTGGTCGTCC TGTTCCACGT CCTGCTTGCT CCCATCACAG CCATCTTCCA	099	CCCARCOCCA ITTICTAAAGA GCAACTITICC AGCAGTACAT ITGAAGGACT TITITAAAAA
CACCCACTTC TATGACAGGC TACAGGACGC GGGCTCTCGC TGGCCCGAGC TCTACCTCTA	720 40	алсаластып даассалалт падалестся псалкитель питичилист всигичис
CTCGAGGGCT GACGAAGTAG TCCTGGCCAG AGACATAGAA CGCATGGTGG AGGCACGCCT	780	AACTAGTITT TAAATITTIAG ATTAGAGGC CACCTACKY AAGTGTACAG TITAATAATTA
GGCACGCCGG GTCCTGGGG GTTCTGTGGA TTTCGTGTCA TCTGCACACG TCAGCCACCT	840	GZTITAITICAA ICANTICZANG ANZOTITADA GIITITICAADA AFTITITITIN ITTAITICAAAA
COGNICACIAC COTACITACI ACACAAGOOT CHOTOTOGAC TICANGOOCA ACTIGOSTOGS	006	
CTGCTGAGGC CATTGCTCCA TCTCACCTCT GCTCCAGAAA TAAATGCCTG ACACCTCCCC	096	GILATISLARI AAARCAACT CTAATGTTTG GCAAAAAAA AAAAAAAAA NTCGAGGGGG
ACHARARAR ARARARAR ACTOGROGGG GGGCCCGGTR COCLAFTCGC CCTRIARAGG	1020 50	GRECOGINGE CANTIDGECE TRANS
	1021	
		(2) INFORMATION FOR SEQ ID NO: 147:
	33	(i) SEQUENCE CHARACTERISTICS:
(2) INFORMATION FOR SEQ ID NO: 146:		(A) LENGTH: 1386 base pairs (B) TYPE: nucleic acid
(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1285 base pairs	09	(C) STRANDELNESS: double . (D) TOPOLOGY: linear

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PCT/US98/04493

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£ SEQUENCE CHARACTERISTICS: (A) LENGTH: 2098 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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S (2) INFORMATION FOR SEQ ID NO: 148:

3 8 AAAAAA GACCTOTTOG COTOGOGOCA CATOGOATOC TTCTAGOAAC ACAGTITGAG AACCACCAAA ATTITIAAAA TGAAAAAAAA GCTOCTCTGA TICAGGGGAT GTOGGICOGG GTAGAACCTO ITICITICAAAA TITITTAAACA TOGGAAGITIC AAACAAATAT AATGTOTGAA ACAGATCAAA 1380 1320 1260 1386

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CICTITICAT GOIGGIGCOT GCICATCITIG CIGAIGCAAA CIAGGAAGIT AGGCIGCAIC

1560 1500

1680 1620

ICOGAGIOGO TITICOCTOGA GAGGIOCTITI OCTOTCICIC AGACTCAGIC ACTOTOTICC

CICCATCIGG TCACIGCAGG IGCCAACCCI ICATCCCCCA IGITITCCTG GGCCATGGAG

GOCTGACCTC COTTTCTGGG GAATGTGGCT GAGCTGTGGT AACCAGCTAC ACCCCAGGTG

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TRACTOTION COMMANDET AGENGACIG GICTIOCICC CAGOTIOCIT COCCOCICOIC

GUAROTOCOT TIGUAGGOOD CIGIOGITICO TOCOGOACOA GIGIOCOTIGG CIGOCATGGO

AAGCTCATCA GOOGCTTOTA CCCTGOTCAC CAAGCATGOT AGCAGCTGCC TGCAFTGTAT

1380 1320 1260 1200 1140

1440

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CCTGGCTGAG ACTGTCAGCC TCCTGGAGGA GTGGGGTCCA CCTTCTTCTT GCCCTATGCA

GIOCAAGCTI CACTICICAC CCAGCAAGGI IGACICAICI GCCICCAIGI CICIGGGCT

ACCITITATES ATSCATOSTIC ACCATAGOOD COTTOCTTOT GACCIOGACE CICCATIGIA

1080

1020

960

900 840 780 720

TOOTOGTGAA COTTOGACCA CAGCATGTCA GTGCTCTAGG GATTGTCTAC TOCAGGGATT AGCACCTICA TCACCAGAGG CITGAAGGAA CCCCGCCATG TGGCAGGGCA CAGGCACTGI 1200 1140

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GANGOCAGOG TOGOCAGCAG TOGOTOGOTO CTTCOGCOCC CCTCCOTOCT CCTTTCCCTG TOCAGOCCOT GOTGAGGAGG GOCGCCTCCC CAGAOTCTGC TCCCCCGTGA ACACAGAGCA 1080 1020

TOTOCCAGGG GOTCCAGCTG GAGATGCAAG GCCCCTGAAG GCGCAGGCTN CCAGNCCGCC

960

ACATEGRACIT GEAGAAGGAC GAGACTETES ANSITSSTEAA COGGETACTS CAGCACATEC 90

33

AMATTERCTT CCTGAMTGAC AMGGTGGAGG AGCGGCGGA NCGCTACATG GACTCCTATG 840 780

30

GAGACTETAT COGGGACETE ACCATGGCCG ATGGGGTTCC TGGTGTGCAG AACATTETCA TOTOTOHGAA CISTGOTIAC TICCAGCAAC TIGAGGGCAA AACCAATOTC ATCCIGCIGG

93 720

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AUGGITTICT CCAGGGATTT AAGGGCCAGC TGATACACCA ATACAACAAG AACAGCTCTG

ACATTECCCT TITEATETTT TETGEGGGCA TIGGTGATAT CETGGAAGAA ATTATECGAE AGATGAAAGT GTTCCACCCC AACATCCACA TOSTGTCTAA CTACATGGAT TTTAATGAAG 8

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CTICTOTOGO TCTACACAAG AAGGTTCCTG TGAGGGCTAT CAGTTGTTGC CTTCTAGCTT GACCTAGAAA TGGGCCTGTC TGGCATTTCA GAGTCAGGCA AAGCAGGCAG GGCCAGGGAG 2

CTTGGCCTAG CCACTGGGCT GGGATCTTCT GGGTCATGTG ACTGTGTATC CAGGAGCAGA

AACTIGINIT CICAGANIC AGGNICIACC CAGCACCAAA GAIGIATITI CAGGAGAACA

CCTTCRATGA AGTGAGACCC TTGGGGAGAA CGGGCTGTGG ATGAAGGAGT GGACTGCAGC CTGGCCATGG TTGCTGAGCA TGGGCAGACC AGTGGAGGCC ACCCTACTGT GTTATCTGCG

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GTCAATCCAG AACTGCCTCT GAGCTCCAGG CTGACCACAG ATCAGCCACA GCCTGATGCC CACTECCTOG GGACTATIGG ATCACTGTCC CCCCACCTGT GTGGCCACAC CATGTGTCCT AGCATGAGCC ATGIGCTICT TIGCCCTICT CIGICCIGIT CCAATCITCT GCCTCCCAGT CTTCTTCCCA CCTCAAGATG AGGICCTCGC CCCCTTGTCT TGGCATAAAA ACACCTTTAA GOCCTICCCT GICCACIGGG CIGGATICAT GITCANACCA CIGGACIGGC AGGGCAACGA TOTOCCATEC TOCCTTOTICS CAACCTCTG CAGGATGCCC TCCCTACCCA MCTYTYCCTG OCTOSTAACT TTOOCOCCTC COCCAAGCCC TOCCAGACTC CCCTOOCTST GATOSCATTC

CIPROCIENC COTTOCCOTO COCTOCTOCT TOCTTOCHCA CAGCAAGCCT

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AGTICCAATIGC AATIGCTICAGG GAGGGATATA AGACCTTICTT CAACACACTIC TACCATAACA COCACAATOT CCTATOTOAG CAGAAGATTO AGAAGTTTCA GATAGCCCAG GTGGTTAGAG 480 420

TOTAL COGGACCOTO ANGUNGANGO TACCTONTAT GOTGENATGE TECHCONANG 360

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TTOCATATAA TOGAAAGCGA TGCCCTTCTT CTTACAATAT TCTOGATAAT AGCAAGATCA TCAGTGAGGA GTGTCGGAAA GAGCTCACAG CGCTCCTTCA CCACTATTAC CCAATTGAGA 300 240 180

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GCAAGGGCGS CGGAGACCGG TTACAGGTGA TTTCTGATTT TRACATGACC TTGAGCAGGT ATGAAGGCCA CGGTCCTGAT GCGGCACCTG GGCGGGTGCA GGAGATCGTG GGCGCCCTCC 120 GOCACGAGOT GOCGCAGGGG TCAGTGOTTC TCTCGGOTCT CGGGACAGGT GAGCACCCTG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

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AGENCETICITE COCCOCCOTTO GGACTICTGAC ATCTTAAGGC TGCACGGTCG TGTCCTTGTC

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

(D) TOPOLOGY: linear

TOGGTGAGGE CATGICTOTG ATCCAAGGIT CCTGGAACTG ACACAGGAAG GGGCTGTGAA

CCCTAAGTGG GTGTMATCTC CTCCRACCGA GGCTTCTMAC CCTGGAGATG GCAGTTACTC

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	373		374
	GCTGGAATTC CTTTGTGGTT TGCTCTTCTG CTTCTCACTC TTGTATTAAG AAGGATTCCA	1740	
	CARACTERICA CHISCAMITTO RETRECTION OFFICEAGACE AGAGITITION GAGGICA	1800	GATGACCOTT CATTAATAAA TTTTGCATCTC ATGCACACCA GTTACTTCCT CTTTGTGATG
4	Charles the rate with the charles to the control to	ע טאמר	STOATAACAA TOTTITISCTA TOCTICITATIC AAGGSCAGAC CTAGCAAATT GCSTCAGAGC
^	GACCTAACC CICCASCICA GCCIGGACA CLIGACCCIA IMANIGADIS GOIIIGCIO		ANTICOTGAAT ITHOTOCCGA GAAGATGGCT ITTOGCTGAAG CCTAATTOCA CAGCTCCTTG
	ACTITIAATIC CIGACACCAG TAAAACCAAA AGGACTICTTG GGGGCTICAGT GTGAGAGCCA	1920	CHIPOSESTY CEACHOLAGE CASTOSTATA PROTECTION OF THE ACTUAL CASTOSTATO
•	GOOTTACCTA CTCTGCCAAG TGAGGACAAA CTGCTAGGCT GTATCCCATA AFTTCAGGAT	1980	
2	GAGAAACATT AACAATAAAA ATTTGTAGTA AACATAACCT CATGANGACT AAAAAAAAA	2040	AICCICIOTO AATACATTAT CITICCATOT TOSSITATIC CAGCCAAAGA CATTICAAGT
	THE SECTION OF THE SE	2098	GCCTGTAACT GATTTGTACA TATTTATAAA AATCTATTCA GAAATTGGTC CAATAATGCA
	AAAAACITGG GAAGAGGCCC UMACCCAII GOOCCIIIIO GOOGGGGGG		COTOCTITISC CCTOSOTIACA GCCAGAGCCC TTCAACCCCA CCTTGGACTT GAGGACCTAC
15		13	CTGATGGGAC GTTTCCACGT GTCTCTAGAG AAGGATTCCT GGATCTAGCT GGTCAGGAGG
	(2) INFORMATION FOR SEQ ID NO: 149:		ATOTITICAC CAGGICACA GCAGCATIGG GICCCICATG GGGITGAAGT TICGITITGGT
20	(i) SEQUENCE CHARACTERISTICS:	20	TCTTGTTTCA GCCCARIATG TAGAGAACAT TTGAAACAGT CTGCACCTTT GATACGGTAT
	(A) LENGTH: 1847 base pairs (B) TYPE: nucleic acid	•	TOCATITICCA AGCCACCAA TOCATITITOT GGATITITATG TGTCTGTGGC TTAATAATCA
	(C) STRANDEDRESS: double		TACTAACHAC AATHATACTT TOTTUTYTYCAT TOTTICATACA CSAAACHTAC CTTTAACHTT
25	(U) LOPULON: Linear	25	
	(AL) SEQUENCE DESCRIPTION: SEL TO SEL TO SEL		
	TOGACCCACG COTCCGAACT GAGGGGGGG CGGGAGCCGG TTGGKGTCTG GTCTTCGGGT	09	TCACATTTTA ATACTACCAA AAAATGGACA AAAAAGTCG AGGGGG
30	CRRCCCCOCC GACCAGACCC TGCCCCCGC GCGGGGAAA GATGGTGCCK AGCGGCCTCG	120 30	
	GECCGCCAC GCGCCGCAC GAGTGAGCCC AGCGCGACCG CCGGCCTCCG CCGAGCAGCT.	180	
	вессеветь вессевеве восемпесе свесевевесь ветранест вителанта	240	(2) INF
35	CHARACTERISTIC CONTROL TO THE TENT OF THE	35	(i) SEQUENC
	ATOTICAGCA TCAACCCCCT GGAGAACCTG AAGGTGTACA TCAGCAGTGG GCCTCCCCTG	300	(A) LENGTH: 1569 base pairs (B) TYPE: mucleic acid
	STORICTICA TRAICAGOST AANOCCCATG GCCATAGCTT TCCTGACCCT GGGCTACTTC	360	
9	ITCABABICA AGGAGAITAA AICCCCAGAA AIGGCAGAGG AITGGAAI'AC ITITICIGCIA	420 40	
	COSTICAATO ATTIGGACTI GIOTOTATCA GAGAATGAAA CCCTCAAGCA TCTCACAAAC	480	(*1) SEQUENCE DESCRIPTION: SEQ ID NO: 150:
	GACACCACAA CTCCGGAAAG TACAATGACC AGCGGGCAGG CCCGAGCTTC CACCCAGTCC	540	
45	CCCAACACT TRABACACTC GRECCEGIG AATATCTCAG TCTCAATCAC CCTAACCCTG	600	GANTIACOCC AACTICOGATIC COCCOGTICOT GAGTICTIGGA COAGTICAAGA AAGCCGTAGC
	COARCIANT BACCOMMISS ADSERBATIVE CTERRACETCA CCCARCIGIA CTCAACCATC	099	CAACGCTOTT CAGCAGGAAG TAAAATCTCT TTGTOGCTTG GAAGCCTCTC AGSTTCCTGC
:	מארכוניאר אוריברוזיסס שמסמיטייסי מסמיטייסי		AGAGGAAGCT CTTTCTGGGG CTGGTGAGCC CTGTGACATC ATGGACAGCA GTGATGAGAT
20	TTAGGGCATC AGATTGGACT TTCAGGCAGG GAAGCCCAGG AGGAGATAAA CATCACCTTC	720	GGATGCCCAG GAGGAAAGCA TCCATGAGAG AACTGTCTCC AGAAAAAAGA AAAGCAAGAG
	ACCCIGCCTA CAGGIGGAG CTCAGATGAC TGCGCCCTCC ACGGTCACTG TGAGCAGGTG	780	ACACAAAGAA GAACTGGACG GGCCTGGAGG AGAAGAGTAT CCCATGGATA TTTGGCTATT
V	GTATTCACAG CCTGCATGAC CCTCACGGCC AGCCCTGGGG TGTTCCCCGT CACTGTACAG	840	
3	CCACCOCACT GTGTTCCTGA CACGTACAGC AACGCCACGC TCTGGTACAA GATCTTCACA	006	
	ACTGCCAGAG ATGCCAACAC AAAATAGGC CAAGATTACA ATCCTTTCTG GTGTTAINAG	096	CIGARISIC ACTIOCACIG CIGCETTIIG GACCASSITG TACCGAAGCA CIACACGCG
9	OSSSCENTIG GAAAAGICTA TCATGCTITA AATCCCAAGC TTACAGTGAT TGTTCCAGAT	09 000	GATECTICCE TECCTITISCS TETECGACEA GAGTEAATGS AGAAGETECG CTGTETECGG

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GOSTGOCTICG ATTICTATATIC TOSTTAGTAA TIGTACATICCT CTTCAGGTTC TAGGGCTCCT CCCATCCCTT GGGGGCAGCC TCGAGTGTAG TCCATTAGTA ATCAGATTCC AGTTTCGACA GOTCOGARAC TOCOCAGTOR ACAGOGTOTO CAROTCATCC TOGACCCAGT OCACAGOGTT CCCACGOOTC COGAAGGATT GACCAGTTAA CCAACATCTT AGCCCCCATG GCTGTTGGCC ATCAGAGGAA CCTTAGAGGC CTGAAATTGT TGCTTCCAGT TTAGCTGCCC CTCAAATTCA GTAGGGAAAA ATTAAACTCT TTGAATCTCC AAACAAGGAA GTTTCAGCAT TCCCTTATGG TITIGGTGCIA TIGAGTITOT TCTTTATTCT TTTATCCCAG TGAAAATIGT TGATCTTGCT ATTITAATIST AAAAATTIIGC ATTITAAAAGG AGTIGGCCCTG TITTICTIGTIST TAAAAACCCCA OTTROCOGRG GGRGARRIOT TGRATICARGA GOGRARACAR CTRCTATGRT TTRTRARCAT COGCTCTTTG ACTOGTOGCA TCCTCAGTAC CCATTCTCCC TGAGAGCGTA GITACTGCTT AGCACGGACA TGCGGCATCA TCGAGTGAGA CTGGTGTTCC AAGATTCCCC TGTCCATGGT AATTICATCI TIATICCGAT IGICAIOGGA AIGATATITA CICIGITTAC TAICAAIGIG TACGAAGATG TICATACCAA ICCAGACCAG GACIGCIGCC TACTGCAGGT CACCACCCTC TATICCATOTO COTOGRATIAC GICCIGCICI GGRAGOTITTA CCAGRAAAACC CCAGCICIAG (2) INFORMATION FOR SEQ ID NO: 151: CTUTGAAAGC TOGTCTTAAA GAAGAGGAAA CTGAATTGAA ACAGCTGAAT TTACACAAAG AGATTATGAC ATTTOGCTCC CCAGTCATCG GCTGTGGCTT TATTTCGGGA TGGAACTTGG AFACTGAGCC AAAACCCCTG GAGGGAACTC ATCTAATGGG TGTGAAAGAC TCTAACATCC E SEQUENCE CHARACTERISTICS: TICCCTICIC CCTITACCCT TCTCCAGAAA TAAAGCAGGT GACAGGGTTT SEQUENCE DESCRIPTION: SEQ ID NO: 151: (A) LENGTH: 1540 base pairs TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear 1560 1380 1320 1260 1200 1140 1080 1569 1500 1440 1020 960 900 840 240 180 120 ઝ 30 23 20 5 50 3 8 2 55 CTCCAAATCC TGAAGCTTTT GGCTTGCTCG TATTGATTTC AGTCTCCTTT GTGGCAATGG GTGTACAGAA CICCATGAAC TATCTICTIG AICTICIGCA TITICATCATG GICATCCIGG TAACTGTGAC AUGGICIAA TICIGCIAAT ATIGICCCGG AAGGAGAGTC AATTACACCT ACCAAGATAC CTGAAATTAC AACTGAAATA TACATGTCTA CTGGAAGCCC CCTGGACTTG TCCGTTTCTC CTTTTGAAGA TATCCGATCA TRAINTICAGO ATTRICAGA CTTTICCTOTT TRAINTITGTG TOTRAINTICT GTATTICATRIC OTCCTOATIOC AAAAGAAGTT AGGAAGGAAA ATCAAGCAAA TACATCTOTT GTTYGAGACA GCCACATTAT GTATITICCGA TITIGCCCAAA ATACTCIGGG AAACAAGCTC TITIGCTIGGG CIGICAGICI GIGGITAGIC ACGIGAATIC AGITAICATI IGACAGATIC CAGAATICCA ATAAATGOCI GOOTGITTIG CICTOTITIT ACCACAGCIG TGCCTIGAGA OTTAACIGT TACTIATGAG GICAATIGGA AAIAAGAACA CCATITTACI GGGICIAGGA TIICAAAIAI (2) INFORMATION FOR SEQ ID NO: 152: ATAATOGATA ACACTGRATT CCCCTATTTC TCATGAGTAG ATACAATCIT ACGTAAAAGA TTGAGGGCCA TOGRAAAAA ATTGGGAAAA GGAAAAACTC AGTTTTAAAT ACGGGAGACT ACTAMAGET TAGCAGCCAT GICTAGCATC ACCITICCIG CIGICAGIGC ACTIGITICA CGAACIGCIG TACACTIGGC ATOSTATOSC TITOSTICAS AACCITOSAT GATSTGGGCT GCTGGGGCAS (i) SEQUENCE CHARACTERISTICS: Ě

GITTAGGAAA CCTAAGICAG CAGAAATTAA CIGGATIAAT TICCCTTAIG

1540 1500 1440 1380 1320 1260

TOCTATION THACTAGATT ATATAGAGCA CATGROCTIA TITTIGHACTG

1200 1140 1080 1020 960 900 840

ACAGTTOCTG CAAGAAAATG TAATTGAATC TGAAAGAGGC ATTATAAATG OCTOTITICA GOCGICATIG CIGCTAGAAT COGICTITIGG ICCTITIGATI

AGACAAGTCC TGAATCTOTG

CCCATAATCT

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TGGTGCAGAA AGATTGTTGG GAACAGACAG GAACCAATGT GGGAATTCAA CTTCAAGTTC

AATCCAGCCA TTCCAGAAAG CACCCCCAGC ACATTAAAGA ATTCCAAATG CTTACTTTTC SCTISISISA TOOGATOTOT STACCATATS TAIGASCOAT TISCISCIOS AATOTOCAAS

AMAMAKACAGT CCCCTAGGTT AMAGAGCAAG TOTACAGGAG GATTGCAGCC TCCCGTTCAG

780 720 660

> CTCAGGGACT GAGTGGGTTC CATCCTCAGT ATTITGATGG GAGCATCAGC TATAACTGGA TIGCTTICCT TTATAIGACT GICCIGGGCT TIGACIGCAT CACCACAGG TACGCCTACA CCTTCCGAGA TGGATGGGTC TCCTACTACA ACCAGCCTGT GTTTCTGGCT GGCATGGGTC

> > 480

ATAATOOGAA CIGTAGCIII TACIIGGCIA COICGAAAAI GIGGIITIGGI ICGGCAGGIC

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ATGAGCTTGA ACATGAGCAA GAGCCTACTT GTGCCTCCCA GATGGCTGAG CCCTTCCGTA

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180 120 8 SEQUENCE DESCRIPTION: SEQ ID NO: 152:

(A) LENGTH: 1719 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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TGRAATTICTC TAAAGTGTAT ATTITIGGGT CCAGTTATAT TATTIAAAA AGTGTTACTT 1560 TGTAAAAATT GTATATAAAG AACTGTATAG TTTACACTGT TTTCATCTTG TGTGTGGTTA 1620 TTGCTTAATG CTTTTTAAAC TTGGAACACT CACTATGGTT AAATAAGGTC TTAAAAGAAA 1680 TGTAAATATT YTGTTAATAA AGTTAAATAT TTTAATGAT

(2) INFORMATION FOR SEQ ID NO: 153:

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120 180 240 30 360

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(1) SEQUENCE CHDRACTERISTICS:
(A) LENGTH: 861 base pairs
(B) TYPE: nucleic acid

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TOCCTTAGAG CAAGGGAAAC ACCTCTCATT CAAAGGAACT AGAAGCCTCT CCCTCAGTGG GOCACGAGGG AAGCCGGGAC GATGTCCGCA TGACAACCGA CGTTGGAGTT TGGAGGTGCT GCTGGGATTT TCCAAGGGAA GCGTTCAGAA TTAGGCNTGT TGTTTCAGGC ATTTCCAAGG GAATAGAATG AATGACCCCA AAATARGGTT TTCTTGGGCG AGGATGTGCT GCATTAGGAA AGSTCACATG ACACAGGCAG AGCAGAGTOG CACCCACCAC AGAATACAGT GTGTGTTATT TAGGGAGACA GCCAGGAGCG GTTTTCTGGG AACTGTGGGG TGTGCCCTTG GGGGCCCGAG TOCTOCTOTC CTATGACCTC TITGTCAATT CCTTCTCAGA ACTGCTCCAA AAGACTCCTG TCATCCAGCT TGTGCTCTTC ATCATCCAGG ATATTGCAGT CCTCTTCAAC ATCATCATCA GGACAGCCGT AAGACTAGGC GATCCTCACT TCTACCAGGA CTCTTTGTGG CTGCGCAAGG TAGRAGOCCA CATTTOCTGC TTTGCAGGGG AGAGTTGGGC OCTATGCATG GGGCAAAACA TIGITACIAG GIGCIGGAGG AACAICCCAG TICACAAAGC CCCCAICTCI ICCTCTGGAG CCAGAGCCTG COGTOGAATC AAGTCCAACT GAAACATCAG AACAAATAAG AGAGAAATAA AAAACAGAAG GAAGATGCTC CAGACCAGTA ACTACAGCCT GGTGCTCTCT CTGCAGTTCC TITICCICAL GIICITCAAC ACCITGGICL ICCAGGCIGG CCIGGICAAC CICCIAITICC ATAAGTICAA AGGACCATC ATCTGACAG CTGTGTACTT TGCCCTCAGC ATCTCCCTTC ATCTCTGGGT CATGAACTTA CGCTGGAAAA ACTCCAACAG CTTCATATGG ACAGATGGAC TICADATGET GITTIGTATIC CAGAGACTAG CAGCAGTGTT GTACTGCTAC TICTATAAAC AGTICATICA AGTICGAAGG TGACCTCTTG TCACACTGAT GGATACTTTT CCTTCCTGGA AACAGCAAAA AAGAATGATT TCTTCTGAAA TTCTGGAACA TGAGGAITCA AGTTTTTAIT ACGAGGAGCC AGCAGTTGAG CCTAAGGTCC TTCTACCTAC CTGGTATTGG CATTTGAGGT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 153: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 154: (A) LENGTH: 1101 base pairs
(B) TYPE: nucleic acid
(C) STRANDELMESS: double
(D) TOPOLOGY: linear (C) STRANDEDNESS: double (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 154: AAGGGAAGG GTTTCCCTNC CCT 2 2 ನ 3 4 2 55 8 23 35 \$

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480 540 909 99 840

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AGTGAGGGTC TIGGGATGAC TIGCIGIGIT CCICAAGCIG CACTITIGGG CCATCICIGC

780 720

PATTITICATI TCAGAAGACT GAAGCAAAGC TGATAGTOTT TGCTGTTTCT TTGGCAGCTA

35 25 6 8 55 80 CACGGICCIC CIGGGIGCCT ACCAAGCIIG GITTIGIACAA AAGCAAGGIG GGAGICTATI CAATTAACCC GITTGAGGCC TAGGITGITT GGCAAGCCCC NGGCCTAAAG TITTAATTCG CTTCATTTAA ATACATCTGT GTGCATACAG ATACGCATAT ATGTGTGTGC GTATGCATAT TAGCOTTTTT ATATATGACC TITGATTICT GITGITIGTA TITTAGCACA GIGTATGCAC GTTTACACTG ATGCCTTCCC TGCCCACCAC AAATTGTGTA CATAGTCTTC AGAATGATAC TICCTAGGGC TOGAAGGTTT AGCAGCAGCC TOGTGCAGTG CCCTOTCATC AAGACAAACC GCAGAGCCAA GGGCCTGAAA GGAAGGGAAA GGGGAGGGTA GCGGGAGGGT AGCAGGTGAG GACACCCIGC ATCCCIGCTA ATAGIGITITG CCACAAGTAT TAGIGAGICT TCCTTATTAA CACCCCTTTC CCCAGCTCCC AACCAAGAGC TOOTTCTAGG CCTOTOTTAT ATGTCATATT TITIGIACATG AGATACATCA CACTTACCTG TGGGCCAGTA TIGIGAAGTG AGTCTGAGTT ATCTCTCATC TGTAGTTTCC AAGAGTTCAG CTGAAGCAGA TGGAGTCCTG CAGCCCAGGA Ě E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 2031 base pa (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear STRANDEDNESS: double LENGTH: 2031 base pairs 660 540 480 420 360 300 240 180 120 8

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INFORMATION FOR SEQ ID NO: 155:

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TGAAAACATG CCACCTCAAG GCTGGGCGCG GTGGCTCACA CCTGTTAATC CCAGCACTTT

GOGAGGCCGA GGCGGGCGGA TCACCGGAGT CGGGGAGTTT GAGACCAGCC TGGACCAACA

20 25 ᅜ 5 S CTCGTGCCGA ATTCCCTGCA G ACTATOTOTO CTTTTTAGAA TAAAGATTAC ATATCATCAT TCCTTTGGGG AAAATTGTTA CAGCCCAGGG CTTGTCTGGA TCAGCACCAA CGATTTTAAA GAAAAAAAGGA AGAGTTTCTT TGTATTTTGT TCTTGAGAGG GGTCAGTCTA GAAGCTAGAT CCTATCAGGA TGAGGAGCAG CHARGETISET GAGTATISCA SCISCATITS CECANAGGGA ATECAGAACA AGTECETECE ATCCAACTIT AATAGTATAC ATTTAANAGA AAAAAAACAA AAGCCCTGGA AGNITGAGGC TOTOTOCTTT CARARGACT TITOTOTOCT AGCTGROTGR CICCITICCIT AGTTCRAGGA GAACTOCICAG CAAGCCTTGA COCCTTATGT ATOTAGCTGA GTCAGCAAGG TACATGATGC TTCAGGTATA AAAACAAGAG AITATAATAA AAAANTAAAA GAACCCTAAA AAAAAAAAAC AGATGAGTAA TIGITAITGA AGATAGICAG IGATAACCAC IGACCAGAIG CTATCAATAC ACAGCIGAGA CAGACCICIO CIGAGIAGCI CIGIGATGAC AAAGCCIIGG IITAACIGAG COGAAACCCT CTACTOCCCC ATAAGCCAGG AAAAGTGAAA AGAGAACACA GTTCCTTTAA CONTIGUENCO TITATTAGTC CCCAAGGCAA ACACAAATAT TAGATTAATA 1101 1080 1020 900 960 840 780 720 660 540 600 480

အ 23 20 5 5 CAGCAGGAAA ATGATTGTAT TIGAGTICCT GIGIGICCAA AACTGAGGCA CCATGTICTT CAGGAAAGGC TACCTAACTT CACATATCTG CAACCAGAGC AGCCACCAAG CATTACTTAG AMATCAMOGO ATCACAMITT CIGITICAGOG GGCAGGAMIA GGCIGIGAMI TOCTAGOACI GOCCCAGGOT GAGTGAGATG AGOTGCAGOT GOCTCATGGO CITOTTAGAG CAGAGAGAGA GICACICTIG CANGGCTICC ANGICIOGIT TOTOGCATTI GOGGATAAGI GCIGAACCAG GOOTAGAAAA TTCACAGTAG GAATGATTOT TAAGAGAGAG TGCTTGGAAC CATGGGTTAA GAGGAAGAAC TTTTAGAAAC CAAATGATCT TAATTGTTAT TGCCCACCCC TGGCTTTTTCC TECEMANIST AGACIAGANI CIGEAGGAIG CEACCEACIG IMIAGITETG CITTECEMGA TITITITIAA GCAATTACIT TITGACITGI TCCTCTGAAA GTGCAAGAGG CGTACACCTT CCAGAAACIG AGGCACAATA GOGTTAIGAC TIGCICAAGA ATAIGTAGCI GCTAGGGGGT AGCATTIGCA GITTOTITGA GGCCICGITG CCAAIGATAG ATCACICCIG TIGACCIGGI AGTATOTCAT TITTACTAAGT TCCTAAACAA ACATTTATGC AGGCAACACT CCTTGCAGAT AUGUCTOCTT GCTTGCTGCT TTTCCTTGCT TTCTCTTGGA AGAGGAAAGG ACTCTGGTCA AGIATIAAGO COCCITITIG CIIGGIGGIA CICIGICIGI GOCIGIGIGI GIGIGIGATA

> 1320 1260 1200

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2) INFORMATION FOR SEQ ID NO: 156:

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TATAATCTCA GCTACTTOGG AGGGYTGAGG CAGGRGAATT GCTTGAACCC RGGANGGCGG

1860 1800 1740 1680 1620 1560 1500 1440

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TIGAGITGAG GATCOTOCCA TIGCACTICC GGGCCTIGGG GCAACAACAG

TOGGRAAAAC COCATOTOTA COTAMAAATA CAAAAATTAGO OGGGOTOGT GGCATGOGOO

CAAAAAYICC GICTICAAMM MRIGCCGAAT TCGATATCAA GCTTATCGAT ACCGICGACC

TOGRAGOGOG GCCCGGTRCC CRATTCGCCC TRIRGNGRTC GTRTTRCRAT C

2031 1980

£ (A) LENGTH: 1981 base pairs
(B) TYPE: nucleic acid
(C) STRANDEINESS: double
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 156:

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	181		382
	COTIGGACCOT GAGCOCTICA COCCTOCGAG TTCCCCCCAG GITGGCTTCC TTCGAFTCCT	09	AATTCTAITG TITTGAACTC TGATTTAAAA TIAAATTGCA GCTGGGCGTG GTGGCTCATG
	TITICITIGGTA TCAACOTITIG ATTGGAAGAA CAACCCCCTC TITIOTCAACC TCAATAATGA	120	CTIGIANICC CAACACTIAG GGAGNAGGR GAATCACTIG ASCYCAGGAG TYCTAGACCA
. 1/3	GCTCACTOTO GAGGAGCAGC TOSGGCACAG CTCACCOTVA TOGTCATTOT TACCCCCAA	180	ATCTGGGCAA MAGAGAGC CCATCTCTT TAAATAAAA GTTAAATTGC TTAAAAAAA
	GACCGCAAAA ACTCTGTGTG GACACAGGAT GGACCCTCAG CCCAGATCCT GCAGCAGCTT	240	
2	GTGGTCCTGG CAGCTGAAGC CCTGCCCATG TTAGAGAAGC AGCTCATGGA TCCCCGGGGA	300	
2	CCTGGGGACA TCAGGACAGT GTTCCGGCCG CCCTTGGACA TTTACGACGT GCTGATTCGC	360	(2) INFORMATION FOR SEQ ID NO: 157:
	CTGTYTCCTC GCCATATCCC GCGCACCGC AGGCTTGTGG ACTCGCCAGY TGCCTCCTTC	420	(1) SEQUENCE CHARACTERISTICS:
15	TECCEGOGCC TOCTCAGCCA GOCGGGGCCC TCATCCCTGA TGCCCCTGCT GGGTNATGAT	480	
	CCTINCTCAGC TCTATCTGAC GCAGCTCAGG GAGGCCTTTG GGGATCTGGC CCTTTTCTTC	540	(C) STRANDEDMESS: double (D) TOPOLOGY: linear
20	TATGACCAGC ATGGTGGAGA GGTGATTGGT GTCCTCTGGA AGCCCACCAG CTTCCAGCCG	600 20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 157:
3	CAGCCCTTCA AGGCCTCCAG CACAAAGGG CGCATGGTGA TGTCTCGAGG TGGGGAGCTA	099	GAATTCOCCA, CCACCCCCCC CATGCCCCTC, CTGCTTTTCGG TGCTGCCTGT ACTGCTGGGC
	GTAATGGTGC CCAATGTTGA AGCAATCCTG GAGGACTTTG CTGTGCTGGG TGAAGGCCTG	720	GOCTITUTICS COCTOSTICS STITUSCCAAG CIVIDOGAGS AGAITOTGGC TCCASTITICS
25	GTGCAGACTG TGGAGGCCCG ANGTGAGAGG TGGACTGTGT GATCCCAGCT CTGGAGCAAG	780	GAGCOGANGA ANOCCCIUTT CONGCAGITT GONGAGGIOT ICCCGCUGAA GGNAITHGGC
	CTGTAGACGG ACAGCAGGAC ATTGGACCTC TAGAGCAAGA TGTCAGTAGG ATGACCTCCA	840	TACCAGCCAG ATCCCCTGAA CTACCAAATA GCTGTGGGCT TTCTGGAACT GCTGGCTGGG
ç	CCCTCCTTGG ACATGAATCC TCCATGAÄGG GCCTGCTGGC TGAACATGCT GAATCATCTC	30	TROCROCROS TCARGOSCCC ACCOARGOTG CAAGARCA GRAACTIGIT CITGATICTO
3	CAACAAAACC CAGCCCCAAC ITTCTCTCTG ATGCTCCAGC ATTGGGGCAG GGGCATGGTG	096	CICATGATOS GOSCIATUTI CACCITIGACA GCICIGAAAG AGICACIAAG CACCIOTATO
	OCCCATISTAG TETECTIGGC CTCACCATC CAGAAGAGA GTGGGAGCCA GETCAGAGAA	1020	CCAGCCAFTG TCTGCCTGGG GITCCTGCTG CTGCTGAATG TCGGCCAGCT CTTAGCCCAG
35	GDAACTGAAC CCAGGAGATC CATCCACCTA TTAGCCCTGG GCCTGGACCT CCCTGGGATT	1080	ACTANGANGG TEGTCAGACC CACTAGGAAG ANGACTICTAN GTACATTICAN GGAATCCTGG
	TCCCACTCCT TTCTTAGTCT TCTTCCAGAA ACAGAGAAGG GGATGTGTGC CTGGGAGAGG	1140	ANOTHGNOCA TCTCTTOTCTC TTTANGCCAT GCAGCTOTCA CAGCAGGAAC ANGGTAGAAC
- 4	CTCTGTCTCC TTCCTGCTGC CAGACCTGT GCCTAGACTT AGCATGCCCT TCACTGCAGT	1200 40	ACAGAGICTA TCATCTTOTT ACCAGTATAA TATCCAGGGT CAGCCAGTGT TGAAAGAGAC
	STCAGGOCTT TABATGGGAC CCAGGGAAAA TSTGGGCCTT CTGAGTCACA TCAGGGACAC	1260	ATTITICION CCICGCACTO CITICICITY TIMOCITIAC TACICITITIO TGAGGAGIAC
	TGAGCAGTGG AAAGGGGCTA TATGTGTATG AATAGACCAC ATTGAAGGAG CACAATGCCC	1320	ATCTTATGCA TATTAACATT CCTCATGTCA TATGAAAATA CAAAATAAGC AGAAAAGAAA
45	TCCIGTOTTIG ATGCCACTTC CCAGGGTGGA GACAGTGGAA AAGAACCGAG GACAGGAAAG	1380	TITABATCAA CCAAAATTCT GATOCCCCAA ATAACCACTT TIAATGCCTT GOTGTAAGTA
	GATTOGGTAG GTGAAGGGGT CAGGGGACTG GTAGTCACCC AATCTTGGAG AGGTGCAAAA	1440	TACCICIGAA CITITITICIG ISCCITIAAA CAGAITATA TITITITIWA AIGAAAAIAA
20	AGCACTOGGG GCTACCOGTT AGCTGCATCT GCCCTGGCTG TTTGCCCGTT CATGTCACAA	1500 50	AACCATATAT CCTATTITAT TICCTCCTTT TAAAACTTA TAAACTATAA HAAAAAAAA
;	ACTECCACTA CTATGTACCT GCAGTGGGGT TGCAGAGATG GGGGAGACTC AAGTCTTACT	1560	алалалала стеса
	CCCCAGAGC TCCCAGGGCC CAAGGAGGAG AATGCTGCCT CCTTTCAGTC TGGTCTACAC	1620	
55	CCACTITICIS GIAGCOICIC IGCITICCIGI AATICIGACI GITITITICCAG ACTCAGCICA	1680	
	AATAGTOCCC CTCCTTAAGC CCATCCCTCG CCCCCAGCCT GAGGTGATCT TTCCCTCCTC	1740	(2) INFORMATION FOR SEQ ID NO: 158:
09	TGAACTATTA GAGCAOTHAC TSTCTSTTCA GTTCGTTTCO CAGGCACACA CAGTGGCATA	1800	(i) SEQUENCE CHARACTERISTICS: (a) LENOTH: 2117 base pairs

χ̈́. SEQUENCE DESCRIPTION: SEQ ID NO:

23 20 2 5 TGAAATGGAT AAGATGCATG CAGGCCTCAT AGATGCCATC AAGCCTTTCC TCGACTATTA COTCAGCAAG ATCATCOCAG AGAATATTTA CGAGGGTOOT CTGAACAGTG ACTATOTCC TOGACAGCAT CTTGCAAAGA AAATCATCTT AAATGCCGTG TTTGGTTTCA TAAACAACCC acrocrocro acaccoroco regrocados gorogadese areageeros gaeroseeeer GITACAGITG TOGATIOGAG GCAACGIGAG TOCCTOTOGG AGGICCATCI TCATATITIGA CCTGTTTGTG GCCACATTGC ACTTTCCACA TGCTTCAAAC ATCACCTTGT ACAAGGATCA COGGUAGAAG COGAGCCTTA GCCGGGAGGC ACTGCAGAAG GATCTGGACG ACAACCTCTT GOCCOGCOTC CTCACCGGCT ACATCTACCC GCGTCTCTAC NARGCCCARG ARACCTCTCA CGCTCTCCCT GCACGGGTGG ACAGGCACCG GCAANARTTT AGAGCGAAGC GAGGGTGGCG COOGTCCGGG CATGAAGCTG GGCCGGGCCG TGCTGGGCCT TOCCTCTTCG CCGAGTGCTG 600 540 480 420 360 300 240 180 120

မ 35 CCCCCTGGAA TACAAACACC TAAAAATGTG TATCCGAGTG GAAATGCAGT CCCGAGGCTF CATCAAGCTC AAAGACATTG AACACGCGTT GTCTGTGTCG GTTTTCAATA ACAAGAACAG TOGOTTOTOG CACAGCAGOT TAATTGACOG GAACOTOATT GATTATTTTG AGCAGAAAGG ATCACAGATG TOGCTTTTGGA TTTTCTTGGAGG AGTGGAAAGC AGAGGGAAGJ TICCCTICCI 900 840 780 720 660

TGAAATTGAT GAAGACATTG TAAGCAGAGT GGCTGAGGAG ATGACATTTT TCCCCAAAGA 1020 960

CAGTCATGAT TOOCAGCCGG TICCACACITI CCACCCCAG TICICAGATA AAGGCIGCAA AACGGIGIIC ACCAAGTIAG AITATIACIA AGTCACTGCC TOGAGTTGGA CICCITICCC TGGAAGAGGA ATCCACTGA ANAGANACAA 1140 1200 1080

TOTTCCTCTT GGGACCACOO GCGAGGACGT TGATOTGACA GGAATTCTCC GAAGCCTCCC GAACACGCAC AGAAGGAAGG CIGGCATIOT TICCACCCC regreccio 1320 1260

TITTAACATT AUGAGAGACT GCTCAGATIC TAAGTIGITG GCCTIGIGIG CIGITICAAG GAAGGAIGAA TAAGITITIAT IGAAAAIGIG GTAACITIAI 1440 1380

TITMAGITCT CATCATTAIT ACATAGACIG TGATGTATCT

TGAGCCCCAAG CACACATOCA TOOCATTTOT TCCACAGGAG GOCATCCCTG GGGATGTGGC

CAGCCCACTC ATCGCAGGGC TCATGATTTT TTACAAATTA TGTTTTAATT

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TOGRGCATGA GCCAGCTCTG TCCCAGGATG GTCCCAGCGG ATGCTGCCAG GGGCAKTGAA

1620 1560 1500

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CCTGTTTTICT AGGTGGAAAG AATCTGATGA GCCTTTTAGG CCTGTTCAGG CAAATTTTGAG TACTCAGGAG ATGGATTICA TICTTTGGCC TCGGAATGAT ATTGAAAAAA TCGTCTGTCT

900 840 780 720 99 600 540 480 420 360 300

GGATGTGGTA CAGCTCCATG CTCCTCGATA TCAGTCTATG AGAAGGGATG TAATTGOCTG

CCTGAAAGGT GTGCGAGATT CCAGCTATTC CTTGGAAAGT TCCCTAGAGC

TTTTACAGAA

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TOACCAGITIT GACTICAAGA TOTATATIGC CITTOTATIC AAGGAGAAGA AGAAAAAAGIC GAAGAAACAA GCTCTGAACA GACTACGTGC TCAGCTTAGA AAGAAAAAAG AATCTCTAGG GGACAATTTIC CAAAGGCTTT AGTACCACCT GTATTTCAAA ATGGGGGACC CAAACTCCCG AAAAAAAKGA CITGITTICIG TIGIACITGA GICITAAGAA AAAGIGGCCC ATAGITIIAGI

AGCACTITIT GAAGTGTCTG AGGTTATIACC AGTCATGACA AATAATTATG AAGAAAATAT

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CATGACIGGA CAGCAGCTIT TOTTICTIGA CCCTOTAATA TGACAGICIG CTAATATIGA

CICITICITY CONTICITY CICCONTY CCCGICTGAC CCCAAACGT ATTGTCCAAA CANAGECTIT ATTITEGEAG TINAGECANN TEIGITITICE AGNANGITAG TINITITETE TOGISTICAGE TOTOTITITAT TOCACACETA AATECTGATT ATAGGETITT CATTITETECG TOTTCCTTAA TCCCTTTTCT AAAAAGGGG GAAAATCCCG ATGGATTTTA GGGATTGGTC

180 120

CAGAAGGIGC AGITITIGGG TIAIAGICGI GATITITCGCI AAICAAICAI

ATTROCAGGA

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SEQUENCE DESCRIPTION: SEQ ID NO: 159:

(C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 2395 base pairs

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SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 159:

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ATAMATTIT GCATGIGCAA AAAAAAAAAA ANAAAAAAAA AAAATCCCGG GGGGGGGCCG

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1980 1920

2117

CTTCTGAGGA TOCTGAGAAC GGTGTCTTTC TTTATAAATG CAAATGGCTA CCGTTTTACA TACTATTATT TOTTACOTIC AATCAGAATC CCCGAAACCT CCTATAAAGC TTAGCTGCCC

GTAACCAATT TGNCCCC

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TIGCCAGICA CICCGITIIG CAAAAGGIGG CCCTICACIG ICCATICCAA ATAGCCCACA

1860 1800 1740

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CACACIGICA ACATTIGITA GAACCAGICT TITGAAAGAA AAGTATTICC AACTIGICAC

AACAGOOTOT COAAGOGTTT TOACOTTAGO AACAATGGGA GOTGTGGGAG TGATTTTGGO

GTOTTTAGGT GAAGGACAAG TAGGTAAGAG GACGCCTTCA GGCACCACAG ATAAGCCTGA

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TRACCIOGIG GATOGGGICI CCTACCAGAA AGCCATOTIC ATAITICICA GCAATGCIOG

COATGATTGA

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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960 1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 1740 0081 1860 1920 1980 2040 2100 2160 2220 2280 2340 2395

ITTCATCATG GIGACTATGA AAAAAGITTI CIGCATGTAC IGAGCGGAA GGACAAGACT GENATICATIC TCANCARICC TANCCAGICA GIGITICICT TCATIGACAG ACAGCACTIG CAGACTCCAA AAAACAAAGC TACAATCTTC AAGTTATGCA GCATCTGCCT CTACCTGCCA CAGGAACAGC TCACCCACTG GGGCAGTTGG CACCATAGAG GRTCACCTCC GTCCTTATAT GCCAGAGTNG AGTACTGACC AGCAAAATGG AGAAGATCAG AGAATGCAGC AGCAGTTTTT TITCHTGITT TCTTACCACT TTAITCTITC AGACITTAAA GAAAATGGAC TCATGCACAG AACACTATGC ATTITGAAAC ITGITCAICC IGGATITITIT IAAAICAITI ITAICICAGA ACTIDADACAA AAATTAGAIG ICGIGCACGG ACTOTGIGAA AGAAGAIGCT ITGCALAITT GCTGCACTGC ATCAGTATCT TACTAAAAT GTGAAATGAA AGGACTATTG TACACTGAAA TECTTABATE TATCTEAAAG CACAAGGTGA TACTCATTTT TATGGTCTTC CCATTTGTGC IGGITITITIGE CICITITISACA ICIGICAICA GIATITIAGAG GGIGAGAAGI GAAIGIAACA GGTATAAATA ACATTITITAA AAACAATAAC TITICCTATAA TCACAGTITGT TCCAGAGCAC IGTCAGATAC ATTCTAATGA CCAGAACTÖG TTTAAAAAA GAAAATACAA CCATGGGAAA GAAATCTTAA ATGAAAAACG CATCTCATTG TAGGCATTTT TGCCTCATAT TTTACTGGGC CATGITITGIT ICCIOGIACT CAIGIAITIT ITITITICCAG AICICITICC CCAAGITICCI ATTGTAAGAG TATTCTGCTG COTGTGGATG CAGTTATACA CATTAAAGCA GATCTGGAGT CTGAAGTAGC TATAAAGCAG CTATAAAACA GAAATACATG CATAGCTGCA GAAACCATGA PAGGIAGAGG ACTITICITY IGGITTIGIY TIGITITIGIY TIGITTIGIY ITIGGITTITA

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(2) INFORMATION FOR SEQ ID NO: 160: 55

GAATTTGAAA AAAAAGAAA AAAAGAAAAA GAAAACCTAA ATAAAATAGG TGAAA

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ITTIACACCA ICCTAAAGAA AAACITIACA AGGGIGITIT GGAGTAGAAA AAAGGITAIA AAGTIGGAAT CITAAAITGI AAAAITAACC AITGAGIGIC AAAGTICTAA AAGCAGAACI CATTITICIOC AATGAACATA AGGAAAGACT ACTGTATAGG TITITITITIT ITCICCITIT AAATGAAGAA AAGCTTTGCT TAAGGGTTGC ATACTTTTAT TGGAGTAAAT CTGAATGATC CTACTCCTTT GGAGTAAAAC TAGTGCTTAC CAGTITCCAA TIGTATTTAG CTTCTGGTTG

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CAGAGAAGAG ATTTTTATTA CAAAGAAAAA AATTCCAGTG AATTGTGCAG AAATGCTGGT

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2120 base pairs
(B) TRPE: nucleic acid
(C) STRANDENESS: double

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1140 720 780 8 8 980 207 1080 COCCOGATAC CACCTGACGT AGTGCCAATC ACACCTCTCG CGTCTCGGCG CCTCGGAGGC TAATGAGGAC GCCTGGGGAA ACGCAGTAAC GGATTTCCGG GTGGACCTTC GCTTTACGGC GETGETGETA AGGGGGTCCC COGCACOCAC CATETISTICC CATCCCOCCC GCCCGAGGCA TIGCAGATIT TOGAAGATOG CAAAGTICAT GACACCOOTO ATCCAGGACA ACCCCTCAGG CTGGGGTCCC TGTGCGGTTC CCGAGCAGTT TCGGGATATG CCCTACCAGC CGTTCAGCAA AGGAGATCGO CTAGGAAAGG TTGCAGACTG GACAGGAGCC ACATACCAAG ATAAGAGGTA GCCAACCTAC ATCAACCACA ATTTCTCCCA GCAGTGCTTG AGAATGGGGA AGGAAAGATA TOTTGOGTAC COTTACCOCA GIOGNAAGCT TOGAGATGAT ATTGACCTTA TIGTCCOTTG TGAGCACGAT GCCGTCATGA CTGGAGCCAA CGGGGAAGTG TCCTTCATCA ACATCAAGAC TCAGCGAGGG GCTGTCATTG CCACGGAGCT GAAGAACAAC AGCTACAAGT TGGCCCGGTG GACCTICCTIOT GCTTTGCTGG CTGGATCTGA GTACCTCAAG CTTGGTTATG TGTCTCGGTA TOSTGAGTTC TTCCCCCCAA CCCAGAGGAA GCGGGAGAGC AGTTTACGAC AGCCCCGGTC GIGITTACGG CGGCCCCCC TGCGCGCCA TGTTTCCTCT TTTCCTGGTT TCTCAAGAGT CACAAATAAG TACTCCTCTC AGTTTGGTGG TGGAAGTCAA TATGCTTATT TCCATGAGGA GGATGAAAGT AGCTTCCAGC TGGTGGATAC AGCGCGCACA CAGAAGACGG CCTACCAGGG GAATCGAATG AGATTTGCCC AGAGGAACCT CCGCAGAGAC AAAGATCGTC GGAACATGTT GCAGTTCAAC CTGCAGATCC TGCCTAAGAG TGCCAAACAG AAAGAGAGAG AACGCATTCG actigcagaaa aagtitccaga aacaattitigg ggttaggcag aaatgggatc agaaatcaca GAAACCCCCA GACTCTTCAG TTGAAGTTCG TAGTGATTGG GAAGTGAAAG AGGAAATCGA TITICCICAG TIGATGAAGA TGCGCTACTT GGAAGTATCA GAGCCACAGG ACATTGAGTG ITICICOCCCC CIAGAATACT ACGACAAAGC CITTGACCCC ATCACCACGA GGAGTGAGAA GCCACTGCGG ASATNCAAGC GCATCTTCCA CACTGTCACC ACCACAGACG ACCCTGTCAT COSCARGOTO GCAAAAACTO AGGGAATGT GTTTGCCACT GATGCCATCO TGGCCACGCT GATGAGCTOT ACCOCTCAG TOTATTCCTG GGATATTOTC GTCCAGAGAG TTGGGTCCAA ACTICITICITIT GACAAGAGAG ACAACTICTGA CITITGACCTC CTGACAGTGA GTGAGACTGC CAATGAGGCC CCTCAAGATG AAGGTAATTC CTTCAATTCA CCCCCCAACC TGGCCATGGA CAACTTCCCC AACCCAAACC COTTTOTGGA GGACGACATG GATAAGAATG AAATCGCCTC ACTICAATIGAG TIGGGATTICCA GGCACTIGTAA TIGGGOTTIGAC TIGGGGTCAGA AGCTIGGACTIC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 160: S 2 2 23 8 30 35 6 5 S 25

TTCAGTCGGC CCTGGATGTC AGCCCAAAGC ATGTGCTGTG CAGAGAGGTC TGTGCTGCAG CCTCCACCCT CACCTGTRAC TCAGGACCAC AGAAGCAAAA GTTCTCACTC AAACTGGATG GGAAGCTGAA GTCCTTCCAG ACCAGGGACA ACCAGGGCAT TCTCTATGAA GCTGCACCCA GGAGTITIGIC CITICCACCGA GACTACGAGG GCCTTTGATG CITAGTGGAA TGTGTGTCTA OGICATOCOT GICTACAGOO TOCOTGATOG CACOTTOAGO TOTGATGAAG ATGAGGAGGA TCACATCTGC ATGAAGCTGG AGGAGGGCAA ATACCTCATC CTCAAGGACC CCAACAAGCA GOCCCTCCC GCCGCRGCGA CCTCCAGAGC CTGGGCTACT GCATGCTGAA GTGGCTCTAC CCAAGGATGG GCGCTTGTTC AATGAGCAGA ACTTCTTCCA GCGGGCCGCC AAGCCTCTGC ACTIOCICIC TGACATITAG CAGAIGAAAT AAAATATATA TCIGITIAGT CITAAAAAAA AATGTGACAG CTGAAAATAT CTTTGTGGAT CCAGAGGACC AGAGTCAGGT GACTTTGGCA GIGGCCIGCC GGCIGCIGGA IGCCCIGGAG ITCCICCAIG AGAAIGAGIA IGTICAIGGA GITTCGGTGT TCACCAGGAC AAATACAGGT TCTTGGTGTT ACCCAGCCTG GGGAGGAGC AMOTEMACAA OTOGAAGAAG CTOTACTEGA ECCEACTOCT GOCCATECET ACCTOCATOG GOCTATIONOT TEOCINTITECS CTATIONNEA AGIOGEANAE ACGIOGECTA COTOGANOSE TACGCCATGC TGAGGAACAA CCTAGAAGCT TTGCTGCAGG ATCTGCGTGT GTCTCCATAT GAGACCCTOC AGAAGTACCT GAAGGTOGTG ATGGCCCTCA CGTATGAGGA GAAGCCGCCC AAGITIGIIG AIAAGCCGGG GCCCIICGIG GGACCCIGCG GICACIGGAI CAGGCCCIC GGGTTTCTGC CATGGACAAA TTGCCTTCCC AAMAMTGAGG ACATCATGAA GCAAAAACAG AGCAGGAGCC CTCACGAGGG GGACCTTGAG TTCATTAGCA TGGACCTGCA CAAGGGATGC 2100 2040 1980 1920 1860 480 360 300 240 180 120 660 90 540 420 900 840 780 720 ઝ 30 25 20 5 3 6 7 55 50 S TCAGCCTCTC TTACTGTACT CTCCGGGAAT GTTAACCTTT CTATTTTCAG CCTGTGCCAC GAACTETATE CTOTTAGGAT CTTCTGAGET TOTTTCCCTG CTGGGTGGGA CAGAGGACAA CTAGGATCCA GGACTGGGTC AAAGCTGCAT GAAACCAGGC CCTGGCAGCA AACCTGGGAA TECETIBEET GIGGGEAGIG GAGAGGETGE TGGGTGTACG CTGCACCTGC CEACTGAGTT CCAGOGIGIT TICCACIAGI CACIACIGIC TICICCIIGI AGCIAATCAA ICAAIAITCI CCCCTCACCA AGGCTGGGAA CAGAGGGGAT GTGGTGAGAG CCAGGTTCCT CTGGCCCTCT GOCAAGGAAC CTTGCTTTTA GCTTCACCAC CAAGGAGAGA GGTTGACATG CCCAGGGCCA CCGCTTCTTT CITGATCCTC TITCCTTAAC CTOTETAGGE AAGETGGETT CECEATTGGE CECTOTGGGT CEACAGEAGE GGCACGAGAT GAGGGGCACC CAGTGCTTCT AGGGCAGGCT GGGTGGTGGT CCCCTAGGTA ATGITAATTA AATGATIGAA ACTIGAAAAA AAAAAAAAAA AAA CAPTICCCTCT GCCCTGAAGC CTGAGTGAAGA CACATGAAGA AAACTGTGTT TCATTTAAAG GTATATATAT ACATATATAT ATTICTTIAA ATTITTIGAGT CITTIGATATO ICTAAAAATC ACCTAGAGTA ANTOGAGAGA CCAMARGCCT CTGATTTTTA ATTTCCATAA AATGTTAGAA TOGOTOGRAGA TOGGRAGAAA COTGRAACTTO TOTTTTCCOTO TOCOTOCOTOC ARCATTROTG 2 ACCACAAGOG AGGGTCTAGA AGAGGCAGCC CTTCTTTGTC CTCTGGGGTA AATGAGCTTG INFORMATION FOR SEQ ID NO: 163: (1) SEQUENCE CHARACTERISTICS Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 162: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (C) STRANDEDNESS: doub (D) TOPOLOGY: linear (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1003 base pairs (A) LENGTH: 2196 base pairs STRANDEDNESS: double AGTGACTICG CCTTGAGTCT ACCTOCCOC 2003 960 660 60 540 480 420 360 8 120

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(2) INFORMATION FOR SEQ ID NO: 161:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 900 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear LENGTH: 900 base pairs 2

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GITTOCCAGO CAGAICAACO IGAGOGIOGA GAATGOCTOG GGCATITITAC GCIOCOICAI CCACOTOMAA GACTOCTOAC GCOACOTOAT COTAGGCACO CAGCAGTTOA AGCCTAATGA

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INFORMATION FOR SEQ ID NO: 162:

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SEQUENCE DESCRIPTION: SEQ ID NO: 161:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 163:

AAGAAGCGGC ACACGGATGT GCAGTTCTAC ACAGAAGTGG GAGAGATAAC CACGGACTTG

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	386		390	
	GGGNAACATC AGCATATGCA TGACCGAGAT GACCTCTATG CTGAGCAGAT GGAAGGAGA	120	TIAITITIAAC ITIGICITIGC AITIGICCIGI AITITAICACA GITICIGITIG AACAGCITITI	1920
	ATGAGGCACA AACTGAAAAC AGCCTTTAAA AATTTCATTG AGAAAGTAGA GGCTCTAACT	180	CAASTATITIG GOGASTITIAT CITISCEATEC TECCETITETS SITTETICA ECCACETISTE	1980
ς.	ANGANGGAAC TOGAATTICA AGTOCCTITT AGGACTTOS GATTIAACOS AGCTCCCTAF	240	CCACTGCAGT TCCTTCCGTG CTCTGTGACT TTAAGAGAAG AAGGGGGGAG GGGTCCCGGA	2040
	AGGAGIACCT GCCTCCTTCA GCCCACTAGT AGTGCGCTGG TAAATGCTAC GGAATGGCCA		TITIANGITY GINGINTY ICICCTIAGC AGNOGACTY GARATTINCA ATTINGGAAG	2,100
10	CCTITITISTOS TRACATITOSA TRAGSTAGAG CTGATCCACT TITRAGOSGST CCAGTITICAC	360	aactaadaga tgaataaact gggttttttt tgttgtttgt ttttgtaada aadaadaaa	2160
	CTGAAGAACT TTGATATGGT AATCGTCTAC AAGGACTACA GCAAGAAAGT GACCATGATC	420	DADADADADA DADADADA DADADADA DADADA	2196
ž	AACOCCATTC CTGTAGCCTC TCTTGACCCC ATCAAGGAAT GGTTGAATTC CTGCGACCTG	480		
3	AAATACACAG AAGGAGTACA GTCCCTCAAC TGGACTAAAA TCATGAAGAC CATTGTTGAT	540	(2) INFORMATION FOR SEQ ID NO: 164:	
	GACCCTGAGG GCTTCTTCGA ACAAGGTGGC TGGTCTTTCC TGGAGCCTGA GGGTGAGGG	009	(1) SEQUENCE CHARACTERISTICS:	
70	AGTGATGCTG AAGAAGGGGA TTCAGAGTCT GAAATTGAAG ATGAGACTTT TAATCCTTCA	660 20		-
	GAAGATGACT ATGAAGAGA AGAGGAGGAC AGTGATGAAG ATTATTCATC AGAAGCAGAA	720		
25	GAGTCAGACT ATTCTAAGGA GTCATTGGGT AGTGAAGAAG AGAGTGGAAA GGATTGGGAT	780 25	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 164:	 .
	GAACTOGRIGO ARGARDCCCO AAAAGOGGAC CGAGAAAGTC GTTACGAIGGA AGAAGAAGAA	840	GCACHGAGTC GGGGGGACGG ACAGGGAGAG GAGGAGAGG GGTCTGCGGG CGGCCGCTAC	
	CAAAGTOGAA GTATGAGCOG GAAGAGGAAG GCATCTOTOC ACAGTTOGGG CCOTGGCTCT	006	CCAGAAGCCA GCAGACGGCA GCACGAGTG GGCTGTCCCC GAGCCAGCC CCGAGGGAGC	120
30	AACCOTGOTT CCAGACACAG CTCTGCACCC CCCAAGAAAA AGAGGAAGTA ACTTCTGAAC	30	CCCCCCCCC CCCCCGNAGG ACGCCCTTVC CAGCCAGCCC GACTYCTAGG AGGAGGGGAG	
	ITTICACCETG AGETECATTE TTECTECAGE CAACCETGA AAATTITIACA TGACATAGAA	1020	GOGGANAAG AGCTCAAGCC TCACCCACGG CCCTGCCCCC AGCCCGGCCA CTCCCAGGCT	 240
35	ACTORATTY TECTTICOTT TICATTICAA GITTIGECAT TIGIGITIAT GGOTTIAGGG	1080 35	CCTCGGGACT CGGGGGGTCC TCCTGGGAGT CTCGGAGGGG ACCGGCTGTG CAGACGCCAT	300
3	GOCCHITICT CTGGACCAAT CTACTCGGGG AAITTCCAGGC CCACCAGGAC ACGTGCCAAT	1140	GENETICOTO CTICATUCC TETICENACET GETGGCCCCC ATGOTECTICS CENGIFICAGE	360
:	OCCCCATTC AGATGCCAAG GGAGGAGGTG TTCTTGAAGA CAGGAGGAGG CTCCCGCTGT	1200	TCANANGGAG ANGGANATGG ACCCTTTTCA TTATGAITHC CNGNCCTGA GGATTGGGGG	- 420
40	TAATAATAT TETTTCATTC TTCTCTTC CTGTCACCTT CTGCCAAGAC ATTGATGGCT	1260 40	ACTIGATOTIC CCTICTICATOR TOGGATICCTC CTTATICCTAA GTCGCAGGTG	
	TCTGACATET TATTTGGTGT CTCAAAGCTG TATTTCCAAG ACAGTGGTAC AAGGTGACCC	1320	CAAGTGCAGT TTCAATCAGA AGCCCCGGGC CCCAGGAGAT GAGGAAGCCC AGGTGGAGAA	540
. 45	THANTHACC GIAICANGGI TCTIGACCAG CACATICAAT CCTCCAACCT ACCCTACTGC	1380 45	CCTCATCACC GCCAATGCAA CAGAGCCCCA GAAAGCAGAG AACTGAAGTG CAGCCATCAG	09
	CATGACCTTC CGCACATCTC TAAGTTTTAT CTTTGCAATA CTCAAGGTTC TCGGAAATTT	1440	GTGGAAGCCT CTGGAACCTG AGGCGGCTGC TTGAACCTTT GGATGCAAAF GTCGATGCTT	9
•	GCTAATGGTT GTGATAAACC ATACAGCTTG AGCCAGTGAG GCAGATTGGG CTGGTGCCTT	1500	ANGANANCES SCENETICAS CAACASCEET TTCCCCAGGA GAACCCAAGA ACTTGTGTGT	720
20	CETCTGAGIT ITCCTGCTTT CCTGCCTCGT GCAGAITCTG AGGIAIAICT GCTGCCTTGG	1560 50	CCCCCACCCT AFCCCTCTA ACACCATTCC TCCACCTGAT GATGCAACTA ACACTTGCCT	780
	AAGACATAAG AAGCAGTGAT ACTCCCTGGC TCGGTTATTT TCTCCATACA ATGCACACAT	1620	CCCCACTGCA GCCTGCGGTC CTGCCCACCT CCCGTGATGT GTGTGTGTG GTGTGTGT	B40
55	GGTACAATGA TAGAAGGCAA AATTGCCACT GTCTTCTTTT TTTTCTCATA TATCTAAGGA	1680 55	GIGACIGICS GIOTITICCTA ACTIGIGACT TIVITICAC TICITICIOS ANGCIANTICS	6
	AGATATATCA GOTTOTIGCCT CATGTACCGC TTCTAGTGAA ATGTAGAGGA AGGCTCAAAG	.1740	CITIVITACI GAACTOTOGA CTCGCTTTCC CAGGCAGGGG CTGAGCCACA TGGCCATCTG	6
	GAGTEAACAT TTAGATETIGG AAGGGACAAG TEATGCETTG GGCCTAGAAT ACCETGATGA	1800	CTCCTCCCTG CCCCCGTGGC CCTCCATCAC CTTCTGCTCC TAGGAGGCTG CTTGTTGCCC	1020
09	GAAAAGAGA GAGGAAGGGA GGCCATATET ACAACANCAN CCTCTCGGCA CTGCTGCTCC	09		

CAGINGINIT GOGACCIOGG AAGGITIGCA GCACITIGIC AICATINITC AIGGACINCI GAGACCAGCC CCCTCCCCTO ATTTAGGGAT GCGTAGGGTA AGAGCACGGG CAGTGGTCTT

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332 6 S 50 8 GENERACION TOCCOPTITOT CONNECEMENT COCCOMINGY CYOGRAGAGE CHARCECCO GATCAGGAGC CAGTGTATAC CGCCCGGCCA CCGCCTTGGT GCCCCTAGAG GAAACGAGAA GOGICGACCC ACGCGICCGG CAGCCGICGI TIGAGICGIII GCIGCCGCIG CCCCCICCG (2) INFORMATION FOR SEQ ID NO: 165: ACACAGIGIT GCIGAAAGGA AAGAAGAGAC GAGAAGCIGI TIGCAICGIC CITICIGAIG E Ě SEQUENCE CHARACTERISTICS: ACAGAAGAAC COTCCCAATC GOTTAATTOT TGATGAAGCC ATCAATGAGC GCGCGCCATG GCTTCTGGAG CCGATTCAAA AGGTGATGAC CTATCAACAG GGTGTCCTTG TCCCAGCCCA AGATGGATGA ATTGCAGTTG TTCCGAGGTG COCYTOCCAC CSCTCOTAGC COTTACCCGC GOOCCGCCAC AGCCGCCGGC SEQUENCE DESCRIPTION: SEQ ID NO: 165: (A) LENGTH: 2933 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double (D) TOPOLOGY: linear 480 420 360 300 240 180 120

20 ઝ 25 2 5 S GOTCATAACO AGAGTOGGAA CTCAACCCAG ATCCCGCCCC TCCTGTCCTC TGTGTTCCCG ACCTICCCIG CITCIGAGAC TICAATCIAC AGCCCAGCIC ATCCAGATGC AGACTACAGI GIGICCCCIG CAPATCTICT CAGCAATAAC ICCAIGGGCT CIGGGACCCT ACCCCTICCA CCCCTTCACA GAGCGCCCGG GGATTCCAGG CCCAGGGCTT CTACTCTGCC CCTGGGGAAT ANGCCCTTCC GIGGITAANT TCITCCCAGG GGCTTCCACG AGGAGTCCCC ATCTGCCCCG TTAGCAACTG GAGATACAAA GCAAGGAGCT GGTGAGCCCA GCGTTGACGT CAGGCAGGCT TICACICCII TAACAAAAAC CITOCIICCI TATCCCACCI GATCCCAGIC TGAAGGICIC алалалала лалалалала стсы TCAACAACAA CAGAAAAAAG GAATAAAATA TCCTTTGTTT CCTAGTGAAA AAAAAAAAA CGAGOTOCOT TOGAGACTCA GCAGGCTCCG TOCAGCCCTT OOGAACAGTO AGAGOTTGAA TGGGCAGCAG AGGCAACTCC CGCATCCTTT GCTCTGCCTG TCRGTGGTCA GAGCGGTGAG ACACCGGGAT GGATGGAGGG AGAGCAGAGG CCTTTGCTTC TCTGCCTACG TCCCCTTAGA COCTOCAATT GGGTCTCTGG CAGGCAATAG TTGAAGGACT CCTGTTCCGT TGGGGCCAGC COGRARCORA CORRACCOTO COCTOTGROC CRITOCTOTI CICIOTATCO IGRICIRIOC 1920 1500 1945 1860 1800 1740 1680 1620 1560 1440 1380 1320 1260 1200

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CAGCICIACO GCGATTIGOT COCTITIGACA GGGAGOTAGA TATIGGAATI CCIGATOCIA

TANAGCAGAG GOCACATOTO ATTOTTATOG CAGCAACCAA CAGACCCAAC AGCATTGACC AAACTEATOG CGAGGTGGAG CGGGGCATTG TATCACAGTT GTTGACCCTC ATGGATGGCC

CAGGACOCTT AGAGATTICTT CAGATCCATA CCAAGAACAT GAAGCTOOCA GATGATOTOG

1440 1380 1320 1260 1200 1140 1080 1020

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20

CICCRACTOT ACCAPATICAG ACTIGNACIOT TOTTICTTCTT GATICARTICOT COTGAGATICA

960 900 840

AGAATGCTCC TGCCATCATC TTCATTGATG AGCTAGATGC CATCGCTCCC AAAAGAGAGA TEAGEAAAIT GOOTOGTGAG TOTGAGAGCA ACCTTOGTAA AGCCTTTGAG GAGGOTGAGA

35 S 50 3 8 8 CTTGAGCCAG AGTAACCCAT CAGCACTGCG GGAAACCGTG GTAGAGGTGC CACAGGTAAC GACCAPTGAT GCCGAGGTCA TGAACTCTCT AGCAGTTACT ATGGATGACT TCCGGTGGGC CTCAGAGGCT GCTCTGCAAG CCATCCGCAA GAAGATGGAT CTCATTGACC TAGAGGATGA ACCTRIGANCA GTAGECHATIG AGACTENCIGG GCATIGTROOF GCTGACTTNG CAGCECTIOTIG TGAGGCCAAT GTCAGAGAAA TCTTTGACAA GGCCCGCCAA GCTGCCCCCT GTGTGCTATT GTTCTATGGA CCTCCTGGCT GTGGGAAAAC TTTGTTGGCC AAAGCCATTG CTAATGAATG TOCTOTOGIAG CACCOLAGACA ANTICOTOMA GITTOGOCATO ACACOTICCA AGGGAGITICT CTOGGANGAC ATCOGGGGCC TAGAGGATOT CANACGTONG CTACAGGAGC TOGTCCAGTA CATECTEANG OCTANCETOC OCANGTECCE AGTITOCCANG GATOTOGACT TOGAGITECT AMATETETTE ATCATTOGCG CTACCAMOOG GOOTGACATO ATTGATOOTG COMTOCTOM GOCTECTEAC CHAGTCATCA ACCAGATOCT GACAGAAATG GATGGCATGT CCACAAAAAA CCAGGCCAAC TICATCICCA TCAAGGGTCC TGAGCTGCTC ACCATGTGGT TIGGGGAGTC ACCTOSCOGE CTIGATICAGO TOATOTAGAE COCACTECT GATGAGAAGE COOGTOTTOG CTITIGATGAG CTOGATTOGA TTOCCAAGGC TCOTOGAGGT AACATTOGAG ATOGTOGTOG 1800 1740 1680 1620 1560 1500 1980 1920 1860 2160 2100 2040 2280 2220

5 5 GCCTAGGGGA TGTCATCAGC ATCCAGCCAT GCCCTGATGT GAAGTACGGC TEAAGCCTCC TAGAGGAATC CTGCTTTACG GACCTCCTCG AACAGGAAAG ACCCTGATTG AAGAGICCTT GAATGAAGTA GGGTATGATG ACATTGGTGG TIGHTGCICC AGACACAGIG GIGGIGGGAT GCGIGCIGIG GAGITICAAAG IGGIGGAAAC TTAAGCCGTA CTTCCTGGAA GCGTATCGAC CCATCCGGAA AGGAGACATT ATGRIGOTIGCC CATTGATGAC ATACTTOTIC TGATGAGAAG ATTCGGATGA ATAGAGITGT TCGGAATAAC AGATAAAGGA GATGGTGGAA CTGCCCCTGA GACATCCTGC ATCCACTOCG AAGGOGAGCC TATCAAACGA GAGGATGAGG ACAGTGGAAG GCATTACTGG TAATCTCTTC CCTCTTTANG GCAATTGGTG CTGCAGGAAG CAGCTAGCTC AGATICCTAGC CCITATIGCA Tricrigico CTTCGTGTAC

> 780 720 660

600 540

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_	WO 98/39448	PCT/US98/04493	WO 98/39448 P
	393		394
			AAAGATTITT AAAAATAGGG CTGTGTTTAA AAAAAAAAC AAAACARGAA AACCAGCAGT
	occimento acimentoci ivividano italciones antalitado accidente	6540	CATTATAGAG AGGTCACACT CTAAGTGGGG TCGCCGCGTG GCCACGCTTC ACGGTCACGC
ν,	CAAGCTGGCC ATCCGTGAAT CCATCGAGAG TGAGATTAGG CGAGAACGAG AGAGGCAGAC	2400 5	TOSTOCOTICC TGCAGTGGCG TOTTTACATG GTCACACGTG TGTGTATCAC CAGTGGGTCA
	AAACCCATCA GCCATOGAGG TAGAAGAA TGATCCAGTG CCTGAGATCC GTCGAGATCA	2460	ACTIGITICAL ATTICITICG TGGCAGTTTG TGTAGACAAT CTTACTGAGC AAAAGGCAAT
	CTITICANGAA CCCATGCCCT TTGCCCCCG TTCTGTCAGT CACAATGACA TTGGGAAGTA	2520 .	_
2	TOAGATOTT GCCCAGACC TTCAGCAGAG TCGGGGCTTT GGCAGCTTCA GATTCCCTTC	2580 10	
	AGGAACCAG GTIGGAGCTG GCCCCAGTCA GGCAGTGGA GGCGGCACAG GTGGCAGTGT	2640	TGACCCAAAA TTGTTTT GTTACTACAG AATACAGAAG ACTGTTTTTTT GCCACACTC
7	ATACACAGAA GACAATGATG ATGACCTGTA TGGCTAAGTG GTGGTGGCCA GCGTGCAGTG	2700	
3	AGCTGGCCTG CCTGGACCTT GTTCCCTGGG GGTGGGGGCG CTTGCCCCAGG AGAGGGACCA	2760	
	GOGGIGGGC CACAGCCTGC TCCATTCTCC AGTCTGAACA GITCAGCTAC AGTCTGACTC	2820	CALCANTACE CALCANAMENT CALCANAMENTAL CONTENTIONS AND MATERIALS
70	TGGACAGGG OTTICTOTTG CAAAAATACA AAACAAAAGC GATAAAATAA AAGGGATTTT	2880 20	
	CATTIGGTAA AAAAAAAAA AAAAAAAAT CCGGGGGGG GCCCGAACCA TTT	2933	AGAGAGGTT CTCCATTCAT TTCAGTCTG CCTGGAGGAA ACTCGGGGAT CATTCTTTC
25		25	AGTIGIGAAG FICCITTICGT GITACACCCT CCACTGAACC CTCAACCTTC GAAATACTCC
	(2) INFORMATION FOR SEQ ID NO: 166:		ACTITICIOG CITICCICAT TITIACTIAL AAATTIACCT TITICIAITT IGCAATITAC
ć		C.	ATCHOTTHGG THTCHTHAA AFTCHCHAA ACHOOCTHGA THAAAAGACT CCHTHHAAAT
00		200	GGAAGCCACC AGTCAGCAGA ATGGAAGCTT AGAGGAACTT GCCTGTGAGC GCTGGTCTTT
	(C) STRANDENESS; double (D) TOPOLOGY: linear		GIGTTIGGTT FIGICATORA ACGARCTITG CTGGGGTTTT TTGCTTTGTT TTGAGGGAAA
35	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 166:	35	TSTCTTGGAG TAAAITTTAA GTICCTGGAG TTAATTTGTT TTACAGGAAT TTTGTTTTT
	TOSGAGAGCC GGCGGGGGANG CGCCTCTCGG CCAGGAAGCG CCTCTTGGAC GCGTGTNACC	09	AAAAAATAG GATCATTCTG AACTTTGGAA TGACCCCTT ATATATTTC TGAAAATGAA
6	GATGCCCAGA AGTGGCCTTG GGCTGGGGAT CACCATAGCT TITCTAGCTA CGCTGATCAC	120 40	aacacitaca tgaaaaaaat ttccaatgaa gatgtcagca ttttatgaaa aaccagaagt
}	GCAGTITICTE GTGTATAATG GTGTCTATCA GTATACATCE CCAGATITICE TCTATATITICS	091	TATTAGATGA AAGCAGGGAG TGAATCTTTA AAACAGACTT GATCACGCAC ACACAATAAG
	TICTIGGCIC CCTIGIAIAI TITICICAGG AGGCGICACG GIGGGGAACA TAGGACGACA	340 .	TCTTTCTCTC CGAAACCGGA AGTAAATCTA TATCTGTTAG AAATAATGTA GCCAAAAGAA
45	GITAGCIATG GOTGITCCTG AAAAGCCCCA TAGIGAITGA GICTICAAAA CCACCGAITC	300 45	TOTABATTTG AGGATTTTTT TOCCARTAGT TTATAGAAA TATATGAACC AAAGTGATTT
	TCAGAGCAAG GAAGATTTTO GAAGAAAATC TGACTOTGGA TTATGACAAA GATTATCTTT	. 09£	GAGTITIGIAA AAATGIAAAA TAGIAIGAAC AAAATTIGCA CTCIACCAGA TITGAACATC
Ş	TITICITIANGT AATCTATITA GAICGGGCTG ACTGTACAAA TGACTCCTGG AAAAAACTCT	420 50	TAGIGAGGIT CACAITCAIA CIAAGITITIC AACAITGIGI ICTITITIGCA ITCAITITIT
3	TCACCTNOTC TAGAATAGGG AGGTGGAGAA TGATGACTTA CCCTGGAAGTC TTCCCTTGAC	480	actitiatia aagotecaaa acc
	TGCCCGCACT GGCGCCTGTC TGTGCCCTGG AGCATTCTGC CCAGGCTACG TGGGTTCAGG	540	
55	CAGGIGGCAG CTTCCCAAGT ATTCGATTTC AFTCATGTGA TTAAAACAAG TTGCCATATT	600 55	(2) INFORMATION FOR SED ID ND: 167:
	TCAAAGCCTT GAACTAAGAC TCAATTACCA ACCGGAGTT TTGTOTCAGT GCCCAAAGGA	099	(1) Cholinger Pubbarmentonton
09	вотносттва тестесттай самсатела статестота атасалтал татттатеса	720 60	

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TANANCTONA GATCATONAG ANGCAGGGCC TCTACCTACA ANAGTONATC TTGCTCATTG GOTGOGNAGO TITANAAITIC COCTTACMOO GOCOCTATAA GOGGAAACOT TOOCGGAATT TCATAGAGCT TTTAAAATGG TTTCATTGGA TATAGGCCTT AAGAAATCAC TATAAAATGC CTTTATGAAT GAACAAAAGC TAAACAGATA TCCAGCTTCT TCTCTTGTGG TTGTTAGATC CYSCTTCTCA CCTCCTGCCA TGATACTGTC AGTTACCTTA GTTAACAAGC TGAATATTTA GCCCGTTACC AAKTCGCCCT AIWGTGADTB GTATIMITAT TITACTAATA ICIGIAGCTA TUAAATTIAA GCATTITICT TITAAAAGAC AAGTGTAATA GACATCTAAA ATTOCACTCO TITIGITITI KOCTIKOGIT ATKGITITITY TCCCITITCT WAGCIATRAG CIGAICATKG ACAAGCTGAG CTGCTGTGAC AGAGGGGAAC AAGATGGCGG CGCCGAAGGG 1140 1080 1020 1620 1560 1500 1440 1320 1260 1200 180 960 900 840 780 720 66 600 540 8 420 360 8 240 120

> 50 25 6 35 30 25 20 5 5 S accendence GAACCCACCA ACCAGGGGCA GGATAGCCTG AAGAAACATC TACACGCAGA AATCAAAGTT GGTACCCUAT TCGCCG AATTATGTGC TGAAAAAAAA AAAAAAAAA AMWMRARASK RRWWACTCGA GGGGGGGGCCC GGTAATTICTG OCATGTCCTC AAAAATGACT CATGACTGTG GATATGAAGA ACTATTGACT CTGATGCTGA TTTGCACTCT GCTGGAATTC TGCCTAGCTG TGCTCACTGC TGTGCTGCGG GATTCACTT CAGTOTGAGT TOGACAAAA TAATATACCA ACAAGAAGTT ATGTTTCTTA CTTTTATCAT CCATTCATAG GACCCTTTTT TITTATCATC TCTGGCTCTC TATCAATCGC CACAGAGAAA CAGCTOGAAC CAACGGGCAC AGTTGGCAAC ACCATCAACT TCTCCCAAGC AGAGAAAACCC GITAACCATT ATAGAAAAGC AAAGCTIGAG TITCCTAAAI GTAAGCTTTT AAAGTAATGA TOGAMACAGO CITACICIGA CITICCCIGGG AGIGTACITI TCCIGCCICA CAGITACATI AGGITRACCA AGCITITIGGI GCATAGCAGC CIGGITIGGAA GCATICIGAG IGCICIGICI TOTOCTTOCT TOTOTOCAAA TITITACOCAA GIGACTIOTA CACTOTIGAA CICTOCTIAC ATTGGGACTA TCCAGATCTT GTGTGGCATG ATGGTATTGA GCTTGGGGAT CATTTTGGCA AGAAACCOTT GATOOGACTG AGAAACCAGA GTTAAAAACCT CTTTGGAGCT TCTGAGGACT TCTTAAGAAA AAAGOGAGAA ATATTAATCA GAAAGTTGAT TCTTATGATA ATATGGAAAA (2) INFORMATION FOR SEQ ID NO: 168: Œ. Ξ ATACCACGGA CTOCTATACA GCCAAAGCCA GTCTGGCTGG AWCTCTCTCT GTTTCATTAT CCTOTCTOTC AMACAGGCCA CCTTAMATCC TGCCTCACTG SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 168 (B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear (A) LENGTH: 945 base pairs

396

AMATGACCIT TAATGACACT ACATITITCAG GAACIGAAAT CATIAAAATI TIATITIGAAT

1816 1800 1740

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INFORMATION FOR SEQ ID NO: 169:

(1) SEQUENCE CHARACTERISTICS:

ACATTAAAAA AAACCATTAT TTCACTGTCA TTTAAAGATA ATGTG

900 840 780 720 660 600 540 480 420 360 300 240 180 120 S

S

(B) TYPE: nucleic acid (A) LENGTH: 902 base pairs

1680

GTAGAAATGA TOCTTCTGCT CAGGAATGGC CCACAAATCT GTAATTTGAA ATTTAGCAGG

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45

CACGCTGTTG GACGCAGTAT AGTTTCCCTC TGAGAAGCTG AGTATCTATG GTGACTTGGA

TACAACICTT GICCICICGG IGAIGGIATT GCTTIGGAIT IGTIGIGCAA CTIGITGCTA

8

AAGCAAAATG TCCNTCAKMT CGSMAATGAG AAATICACAA GCGCACAGGA ATTTTCTTSA AGATGGAGAA AGTGATGGCT TTTTAAGATG CCTCTCTCTT AACTCTGGGT GGATTTTAAC

TCCCAGGTAC GCACCACATT TGGAGCCAGG AGCCCTACCA AATTTGRGRG RAMCMTCTCT

႘

GACTITITAT CTICAAGCCG ATGACGGAAA AATAGITATA TICCRGTCTA AGCCCAGRAA GOTGAGGTCA' TICTIGGAGTG ACATGATIGGA CTCCGCACAG AGCITCATAA CCTCTTCATG

30

GAGACAAGAA CAACITATGT CCCTGATGCC AAAAATGCAC CTACTCTTTC CTCTAACTCT ATCTGATGAG CAATATOCTT GCCATCTTOG KTGCCAGAAT CAGCTGCCAT TCGCTGAACT

AATTGACTTA AATCGAACTA AATTGGAATG TGAATCTGCA TGTACAGAAG CATATTCCCA

25

GICTIGCCAC COGGCCIGIC AGTIGACCIA CCCCITGCAC ACCIACCCIA AGGAAGAAGA CTICTACCCA TOTCAGAGAG CTICCAGOCT CTITTCAATT TCTCAGTTTG TCGATGATGG

20

GOCCOGAGOT TCGGGGAACCG CTTCGGCTGA AGCATTTGAC TCGGTCTTGG GTGATACGGC

5 GAGCCTCTGG GTGAGGACCC AACTGGGGCT CCCGCCGCTG CTGCTGGTGA CCATGGCCTT

GAAAACAGCA GENACOTICO GTGGGGGGA GAAGGGGGCT GGCCCCAGGA GGAGGAGGAA ACCCTTCCGA

GOCAGGAGGC GGCAGTTTICT GGCGGGTGAG GGCGGAGCTG AAGTGACAGC GGAGGCGGAA

10

TITCGGGTCGA CCCACGCGTC CGGCCAGCCT AGGAGAGAA GTTCGTAGTC CCAGAGGTGA

SEQUENCE DESCRIPTION: SEQ ID NO: 167: (C) STRANDEINESS: double (D) TOPOLOGY: linear

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		398
(C) STRANDEONESS: double		TECTICETITIG TECTRACETE TECCEATETS GALCETEANT CECCAGETIT CECACTITICA
(U) TOTOLOGO : THEORY (U) (U) (U) (U)		GCAGICCITI GCICICITIG CITCIACCIC AAAIAGCCCC AGGAGIGGC TITIAGICICC
(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 103:		5 AATATGGAGG ATYTCAAGGT TCTCCTGGGG GATGGGGATT GGGATGGGCA GAATCTGTTT
	00 ;	TOSANCTICCG GOTTANTICC AGTIGOGICIA AAADCAGAG TGGGCCTTTC CCTCTCTTAT .
otchcadac tecgactec aactaaatta actgaggaa agtatgagct gaaagaggg		COCTIGAGGGT GOSTAAGAAG GACTGTATICT ACACCTGTTIC ITCCCTACCT TCTCTTTTIGT
CAGACCCIGG ATGTGAAATG TGACTACAGG CTAGAGAAGT TTGCCAGCAG CCAGAAAGCT	180	THEGENOSCC TCATTCTANG TTCCTCANGN GAGTCCTTGG CTTANAGCTG TAGCANGGST
TOSCAGATAA TAAGGGACGG AGAGATOCCC AAGACCCTGG CATOCACAGA GAGGCCTTCA	240	STOCTAGGTG GOGGATITIGG AGCAAAACCG TCGAGTAGGC ATGATACTGG TATGGAGTGG
ANGANITICCE ATCCAGTCCA AGTGGGGAGG ATCATACTAG AAGACTACCA TGATCATGGT	300	GCCTGCAAAA TCAGACAGAA ATGGCTTGAG AAGCCGCAGG GGAGCATGCC TGTCTCTCAG
TTACTOCOCO TCCGAATGOT CAACCTTCAA GTGGAAGATT CTGGACTOTA TCAGTGTGT	360	TGATAGAGTA TOGGAGGAG CTCCCTAGCT TGGAAAATGA GAATTGAAGG GOTTATGAAC
ATCTACCAGC CTCCCAAGGA GCCTCACATG CTGTTCGATC GCATCCGCTT GGTGGTGACC	420	AAATAGGATG CCTAGTTGAG GATGTTCCCA AAGTTTTGTC CAATCTTATC ATTAGTAGAT
AAGOGITITI CAGGGACCCC TGGCTCCAAT GAGAATTCTA CCCAGAATGT GTATAAGATT	480 20	TITIATARACCO ACAGAGAA ACCAGAAAGG GAATAATIGIT ACTITIGGANG CITITATITIT
CCTCCTACCA CCACTAAGGC CTTGTGCCCA CTCTATACCA GCCCCAGAAC TGTGACCCAA	540	TITOTICTMOS TOTOSCITITG TACATOCACA AGAATOCIAT ATGCTGCACA TITITOCCTTT
GCTCCACCCA AGTCAACTGC CGATGTCTCC ACTCCTGACT CTGAAATCAA CCTTAGAAAT	600	
STGACAGATA TCATCAGGOT TCCGGTGTTC AACATTGTCA TTCTCCTGGC TGGTGGATTC	099	
CTGAGTAAGA GCCTGGTCTT CTCTGTCCTG TTTGCTGTCA CGCTGAGGTC ATTTGTACCC	720	CORRESPONDED CHECKER CHICARTERIC CYCRITISTIC CRICALITY CONCERNO CHICARTERIC
TAGGCCCACG AACCCACGAG AATGTCCTCT GACTTCCAGC CACATCCATC TGGCAGTTGT	780 30	
OCCANGOSAG GAGGGAGGAG GTANANGCA GOGAGTTAAT AACATGAATT AAATCTGTAA	840	TAILAGETAT GGLCAACEGG GITTCAICIG TAILCICIC TITTCALCIG TAITGITTAI
TCACCROCTA AAAAAAAAA AAAAAAACH CGANCCTNGG TTTTCAGCTC CATCAGCTCC	006	TGAAANTCCA AGACACTATG CCAATGCAAC COTGACTACT TTOGGAGATT GGTAGTCTCT
Ę	35	TITIGATIGATS ATACTICATES GOTICACTAT CATABICACA TCAGOTICTIC TTITITICATIT
	705	Taatisttaac Taatisaasit Ccagagatos gccttagaaa tstottttaa gaattaagaa
	40	GGAGTCTCAA AAAGAAATGA GAGGGATGCT TCCTTTCCCC TTGCATCTAC AAAACAAGAG
(2) INFORMATION FOR SEQ ID NO: 170:		AGAGACTIGTT CTGTTGTAAA ACTCTTTCAA AAATTCTGAT ATGGTAAGGT ACTTGAGACC
(1) SEQUENCE CHARACTERISTICS:		CITCACCAGA ATGTCAATCT TITTITCTGT GTAACATGGA AACTTGTGTG ACCATTAGGA
(A) LENGTH: 1885 Dates (B) TYPE: nucleic acid	45	TIGITATICAG CITICIACTIGS TCTCATAACT CTGGTTTTGG AAGAATAATT TGGAAATTGT
(C) STRANDEINESS: double (D) TOPOLOGY: linear		TOCTOTOTIC TOTGAAAATA ACCTCCCCAA AATAATTAGF AACTGGTTGF TCTACTTGGF
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 170:	S	AATTIGACAC CCIGITAATA AGGCAATTAT TICTGIGTIC ITAAACAGTA TAAATAGTIG
AGAAAACAAC TGAAAAACCA CATTTTTCTA CATACAGCTG GGGAGGTAGC TGAGAACTTG	9	TAAGTTIGCA TGCATGATGG AAAAATAAAA ACCTGTATCT CTGTTAAAAA AAAAAAAAA
GCACTGCGCA CACATACTAG GTTGAAAGAG AGTTGAGGAA ACCAGAAGGC CAAGTGGATC	120	алалалала лалалалала лал
TOCTOSCAAA CCCTGAACCT GTCTCCTGCG CTTGCTCTAC AGTTCTGAAG TTGAAAATCC	180	
THITCATGCC TAGGATCTGC TISAGITATA AACCCCAAGG CAGCCATGTC ATAGACTAGT	240	
CTICLEAGUE CONTINUES CONCORRAGA CONSTRACA CONSTRUCTOR	300	(2) INFORMATION FOR SEQ ID NO: 171:
	09	(1) SEQUENCE CHARACTERISTICS:

45.

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v	(A) LENOTH: 2100 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 171:	
10	TACATIATOTA TOTACACACO OTACCCAGAG TEOTACTOTO GCAGCOTTCA AAAACATACC 1	120
i	ATCAGAAAGA GTAGOTGCTG AGATAAGGAA ACTTTGCCAA ATGABAGAAA GTCACTCACT 1 TCCAATATCC CCTCTTCAAG CGGCTACCGT GRAAGGGCT GCAAACACAT TCCCTGAGCA 2	180
15	TOCCTITOCIG ATACAGCITC TITATATITA TATCCTACTG GATOGTAGCA TATTGCTAAG GTTTCCTGTA CTCTGCTTCA AGOGRATGTA AGYTTTATGG CATTGAAACA TTTAGGAAAA 3	300
20	AAAAAGATOT TTAAGAGAAT TAATAGAGCC GTAGTCTGTA TTAGGATGTG TGTCATATGT 420	22
X	GROTTETARIA AACTAAGCAR COGROGOTET AGAGROTENA AGROTCAGCA CATTCCTTCT 480 CCCTTTTOSCT CTCAGGCTAA CATGAGAGAA AARAGAAAAAG TCTTGGCTGT GGGGATTGGA 540	480 540
5	AGCICAGGG GCCAAATGIC CITGCCAGAI CCTIAGAGGA TIACITIGAC ICCTAAAAAT 600	. 5
30	ТОЗЛАСЛАСА ОСАТНОССТА САОТОЛТОСА ТТСТТАСССА САООТОЗСАА	No.
	TAGGAGAGGG TOCHTGTAAA TAGGACGAGG TAGACAGTGC ATGATTGTAG GAGAAGGGTT 780	oō.
بر	GAAGGGAGGA CATGATICCA AAAAAGAICG TICTCAATGI GICGICTGAC ICAACCAGCI 840	-
í	GGCAGATTAC ACTTOCCAAG TCGTTCCCTT TCCTTCTAAG TCAGTTGGCT CCATATTCAC 900	ō
	тпалатилос спстотитов осилловама аписетские тпилестити теспловама 960	ō
40	CTCTTGGTGT CCTCTTGGTC ATAAAGTTGT CTCCTACCTA ACCCAGTTTT ACCAAATGGA 1020	ຄ
	AGTIVARAGGG GACAVACTAT GGAAGATGGA CTCCATGCCA TTGCAGTCAG CCACCATTCT 1080	- ■
7	CTTTTCCATA TAAGGAGCCC CATTACATAA GCTACGGGTG AGGTTGGAAC AGCTATGTTT 1140	-
ţ	CATAATTICA AGAGIGIGAC CACCCIGCIC TAGICAICAT CATIGGATGA AICCAGITGA 1200	õ
	CTCTTTGGCA AAAGGGTGAT ACTTTTCACT AAAAATGCCT ACTCTTCCTG TTGATGTTCC 1260	6
50	TITICIGITI TIACCITGIC CAATTICCAC ACTAGICATI TITITIAITI TITIAGAGGAT 1320	100
	CAGATITTAG COCTOGAAAA TGAGTICAAA AAFITICAGIG TAATGICATA AGGATGITGG 1380	ä
V.	GATACAGAGA TITITITITIT CCTTGGAAAC AAATGGACTG GGAAGAAACA CAGCATGGCT 1440	-
۶	THECHCICAR TETCARICIG AUGAITARGA CONTEGRAGA TAGICTIANE TARAGETTAA 1500	5
	АПОГПОТИТА СААСПОСАТА САПАСОССО АСАПОСТСКА ААССССССТИ ТЕСТСТАПОС 1560	9.
8	TARATOTORE TACTARGROE AGENCTITECT ACTROCTARG CREARTERTA GEOGGEOCT 16	620

2 5 TATAAGCAAG TGATTIAGGI ATTTTCTTTT GIGTTTAIGC ATTAICIGAC TATATTAAAA CACTGATOTA TCCAAAATAG CACACATAGT TCAGTATGAA AATAAGAGAA TAAAATCTGT GOTTICTCTT TITTICTTGTA TICTTAGCAA ATTGCATTTA TICACTACAT TACAAACCAT THIRITIAGA AATOCAATAA ACTICTTATT ACATITAAAA AAAAAAAAA AAAACTOGAA ATAAGAATGA ATATGGAAAT TATATTTCTT TTTTCTGTAA AAGAGTTGCA ACTACTTTAT CCIGITITIC TATITACCIT CTATCAGITT TCTCTACCAA TTATOTTTTT TCAATGCTCT AAGAATATOT CIATTITICAT AIGIGIGATA CIGACAGAGC CATGGTATTC CTAAAATATA GATGAGCTGC TAGTCTGAAT AACATTCCCT GACTTAGGGA AAGGCACACA AAAACATATA 2100 1740 1680 2040 1980 1920 1860 1800

(2) INFORMATION FOR SEQ ID NO: 172: (1) SEQUENCE CHARACTERISTICS:

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 172: (A) LENGTH: 1930 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear

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8 3 6 35 55 30 GOTOCTOOTO GTOGGGGGG GCGGCATCGG CTGCGAGCTC CTCAAGAATC TCGTGCTCAC CCTTTGANTG TOGTCCCGGG TGCNGATTGG CAGCGCCTCC GCCGCGGCTC GTGGTTGTCC THANGATGAC ATCHOGTATC TOTTGACHAT GGACHAACTA TOGCOGHHAA GGAAHCCTCC AACGGAAGCC GAAGCCAGAG CTAGAGCATC TAATGAAGAT GGTGACATTA AACGTATTTC AGCTOCCCGA AACCATOTTA ATAGAATOTO CCTGGCAGCT GATOTTCCTC TTATTGAAAG TGACTATAAT GTGGAATTTT TCCGACAGTT TATACTCGTT ATGAATGCTT TAGATAACAG TOTACTOCAG TITTACCCGA AMGCTMATAT COTTOCCTAC CATGACAGCA TCATGAACCC ACAGITITIG TITCAMAAGA AACAIGITGG AAGAICAMAG GCACAGGITG CCAAGGAAAG COSTITICIOS CACATOGASOS TGATIGATOT GGATACTATI GATGIAAGCA ACCICAACAG COCCATOGCA CTOTCOCOGO GOCTOCCCCO GGAGCTOGCT GAGGCOGTOG CCGGGGGCCC TACTRAGGRA TOGGCTRAAT CAACTGGATA TGATCCAGTT AAACTTTTTA CCAAGCTTTT AGAAGATGCT GATCAAGAAG TATCTCCTGA CAGAGCTGAC CCTGAAGCTG CCTGGGAACC TICAGAACCT ATACATTICCA TOGTTIGGGC AAAGTACTIG TICAACCAGT TGITITGGGGA TURGITOTOAT COTARGOOGA COCRGAGARA CITTOCTOGO TOTACARITO GTARCACACO TOGRANCAGOT GOSTATOTTO GRICARGIARO TROTATORARA ARGOSTOTICA COCROTOTTA 540 300 240 180 900 780 720 660 60 480 420 360 120 840 60

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	. 401	
	AGITECOTIO GACTOGOCTO AAGIACAAAG TCAAGGAGAA GAAACGAATG CATCAGATCA	096
	ACAGAATGAA CCCCAGTTAG GCCTGAAAGA CCAGCAGGTT CTAGATGTAA AGAGCTATGC	1020
5	ACOTETITIT TEAAAGAGEA TEGAGACTIT GAGAGTICAT TIACEAGAAA AGGGGGATGG	1080
	AGCTGAGCTC ATAIGGGATA AGGATGACCC ATCTGCAATG GATTTIGTCA CCTCTGCTGC	1140
2	ANACCTCAGG ATGCATATTT TCAGTATGAA TATGAAGAGT AGATTTGATA TCAAATCAAT	1200
:	GOCACOGRAC ATTATTCCTG CTATTCCTAC TACTAATGCA GTAATTGCTG GOTTGATAGT	1260
	attogaagga ttogagattt tatcaggaaa aatagaccag tscagaacaa ttittitgaa	1320
15	TANACANCCA ANDCCAAGAA AGAAGCTTCT TOTOCCTTOT GCACTGGATC CTCCCAACCC	1380
	CANTIGITAT GIATGIGCCA GCAAGCCAGA GOTGACIOTIG CGGCIGAATG TCCATAAAGT	1440
20	GACTOTICIC ACCTIACAAG ACAAGAIAGI GAAAGAAAA ITIGCIAIGG IAGCACCAGA	1500
1	TOTCCAAATT GAAGATOGGA AAGGAACAAT CCTAATATCT TOCGAAGAGG GAGAGAGGA	1560
	AGCTAATAAT CACAAGAAGT TGTCAGAATT TGGAATTAGA ANTGGCAGCC GGCTTCAAGC	1620
25	AGATGACTIC CTCCAGGACT ATACTITATT GATCAACATC CTTCATAGTG AAGACCTAGG	1680
	AAAGAACGTT GAATTTGAAG TTGTTGGTGA TGCCCCGGAA AAAGTGGGGG CCAAACAAGC	1740
30	TGAAGATGCT GCCAAAAGCA TAACCAATGG GCAGTGATGA TGGGAGCTTC AGCCCTCCAC	1800
3	CTYCACAGCT TCAAGGAGGC AAGATGGACG TYTCYCATAG TTGATYCGGR TGAAGAAGRT	1860
	TCTCCAATAA TTGCCCGACG TTCATTGAAG GAAGGAGGAG GAGGCCCGCC AAGAGGGGAA	1920
35	TTTAGGANTIG	1930
40	(2) INFORMATION FOR SEQ ID NO: 173;	
45.	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1509 base pairs (B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 173;	
20	GOCCTGGCC TCTGGGCTGA GCCTTGCTAG GCACTCGGGG TGGCTCTAAG GGGCAGGAT	09
	AGGCTGGGG AGGCCGGCC TGTGGCCCTG ACCAGCCCCT TCTCGTGCRG GTTCCACCC	120
55	GATECAGGTG GTCACGGTGCT TGACGCGGGA CAGCTACTTG ACGCACTGCT TCCTCCAGCA	180
	CCTCATEGIC GTGCTGCT CTCTGGAAGG CACGCCCTCG CCGGAGCCTG TTGACAAGGA	240
	CTICIACTICC GAGITITOGGA ACAAGACCAC AGGGAAGATG GAGAACTACG AGCTGATCCA	300
99	CTCTAGTCGC GTCAAGTTTA CCTACCCCAG TGAGGAGGAG ATTGGGGACC TGACGTTCAC	360

420 480 540 8 9 720 780 840 900 960 020 1,200 1260 3 20 TOTOGCCCAA AAGATGGCTG AGCCAGAAA GGCCCCAGCC CTCAGCATCC TGCTGTAGGT GCAGGCCTTC CAGGTGGGCA TGCCACCCCC TGGGTGCTGC AGGGGCCCCC TGCGCCCCAA GACACTECTG CTCACCAGCT CCGAGATCTT CCTCCTGGAT GAGGACTGTG TCCACTACCC ACTIGOCOGAG TITIGOCANAG AGCOGOCOGA GAGAGACAGO TACOGOCTIOS ACGATIGOCOG COGCOTICOS GACTIGACO GAGTOCICAT GGGCTACCAS ACCTACCOGO AGCCTICACO GOGGRAGOTOC CAGOTOGCCC GOCTAGAGCC AGCCAGOGCC OTGAAGTCCA GTGGCAGOTO CICOTCITICO ATCACOTOCA AGGICATGAC CICATOGGCA GIGTCACCCT GGACCACTIT TITIGICCCCA GIGCIGAGAG CAGAGAGAAG CICAICICGC IGITIGOCICG CCAGIGGGAG OCCUPATORS GOOTSAACT GOOTSTOAD CTCACCOSOT ACCOCAGGO ACAGOCAACO TISTOSTOTOC AGCOTANCEC CTACTIGGGG AGGOLADCAG GCTTTTTGTGT TOTOCTAAAA TOTTITATIC TECETITIOG ACCITIANTIT CACTOTECTE GEAGAGATG TGAACATOTG TOTATION OF ANALYTIC TOTAL GENERALA TOCCOSOCC CTCAGGGTG traditation atchaectice cheagostast henaecostae acheeratat estatetaen GTITOTGGGAC COTTOTTAAC ACOTGACACT GTGGGTCTGA CTTTCTCTTC TACACGTCCT GGCATCTTGC TGCTAATCCT GAGGCTGGTA GCAGAATGCA CATTGGAAGC TCCCACCCCA Trangchgag Titgacaccg Tchaarara aaaaaaaa aaaaaaaaa aithetgogg TICCIGAAGT GICGAGICCA GICCITIGIT GCTGITGCIG ITGCIGTIGC IGTIGCIGIT TATTOTICIT CAAAGTGGAG GTCTCCCCTG ATCCAGACAA GTGGGAGAGC CCGTGGGGGC AGGGALCTIG GAGTIGCCAG CACCAAGCGT GATTCCTGCT GCCTGTATTC TCTATTCCAA TOGACCCCAS GOSTCOGIGG TITTCCACAG ANGOTTAGAC CCTGAAAGAG AIGGCITCAGC ACCACCTATG GATCTTGCTC CTTTGCCTGC AAACCTGGCC GGAAGCAGCT GGAAAAGACT CAGADATICIT CACAGIGAAT GGGATTCTGG GAGAGTCAGT CACTITCCCT GTAAATATCC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 174: (A) LENGTH: 3173 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 174: CCTCAAGGG 2 2 ន 25 റ്റ 35 **4** 45 S 55 8

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8 S TCCAAGAAAT CAAAAGAGCC ATCCAAGCTA AGGACACCTT CCCAAATGTC ACTATCCTGT ACAACCACOO TCTCAOGAGA TOTCTGATTT CCACAGACAT GCACCATATA GAAGAGAGTT TRANSTINCA STOTSTITICS STITIGGSTCS TOOSTACAAT ACTGATATTG TOCTOASTAG AMATTEGGEA CAGETGAGAG GAGACACAAG GAGCAGEEEG CAAGCACEAA GTGAGAGGCA 240 180 120 69

5 Ê SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 175: (A) LENGTH: 991 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

3 2 INFORMATION FOR SEQ ID NO: 175:

6 35 30 25 20 CAGTCAGTAT GIGIGAAGAT CCCIGGIGCG TGGCCTTCAC CACGCATCIT GAGCAAATTA GCAGGGTCTA GAGACTOCTG GGACACTTTT CTTGGAGTGC TACTTCAGAA GCCTTATAGG TAATATTTT ATGCCACACT GGGATAAACA AGCAAGATTG CICACTTCTG GAAGCTGCAI OCCTANATOG ACADATOGAT GENTACCCTT CCTGANATGA CTCCCTTCTG ANTGANTGAC AGCCATGCCC AGTATTCCCA CTCTCCAAAA GGAACTGACC AGCTTATATT TCTCACACTT GCATAACAGG CTTAGTAAGT CCAAACACAG ATGACAGTGC TGTGTGGGTC TCTGTCAGAG AGCAGAATGT GGGCTGCATA TAAGCACACT CATCCCTTTG TCTGGGAATC TTTGTGCAGG GGAAAATOTA CCCTTCGCTT GAGGCAGATG CAGCCCTTCC CCCGAGTOCA TGGCTTGGAG GGATCAATAT TTTGCACACC TGTAATAGGC CATGGCACAC CAGCCAAGAT GCTCTGCTCA ATGACTAGAG AAAGCAGGTT ACCTAGTATA GTTTTCCCAA ACTTCTTCCC ATCATAGCAC ATGTAGAAAA CAMACTOCCA CATCTCAGCC TGTAAGCAMA GCAGGAMACC TTCTGCTGGG CATAGCTTGT CCANTACTOG CAGGITCCCT GGATCCAGAT CITICICITGCC CAACTCITAC TOOGAGATIG ATGAAATTGT GATCTAGGCT GCTGGGCTGA ATTCTCCCTC TGGAAACTGA GTTACAACG COGGSTIGGT GAGCGIGGTG GCTAIGTICT TICTGCTTGT TCTCATTCTG TCTTCAGTGT TIGIOGCICI CAGCCATGTA GACACACTOT CCAAATGGAG IGTIGGAAAA IGTICTITICI TIOCIGATAA GATOOOGAAA GCCAGCACAC AGGACAGTAA ACCICCIGGO ACTICAAGCI AGTCCAAGGT GCTTCCCTCC AAGGAAGAGC CAGTGAACAC AGTTTATTCC GAAGTGCAGT ATTITICITIC TGGCCAAGAT TICCTICIGI ATCACICCAA GCAGCCICAG CAGAAGAAC ATATCATOGC TICAAOGAAC ACCCAGCCAG CAGAGICCAG AAICTAIGAT GAAAICCIOC TITITOTICCO TITOTICAAG AGAAGACAAG ATGCTGCCTC AAAGAAAACC ATATACACAT GCCTCTTGTG ACTGGAGGTA ACAACCCTGC CCAGTAACTG TGGGAGAAGG 1140 1980 1920 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320 1260 1200 1080

> 8 35 မ 23 20 5 5 S GAMOCCKOCT GOOGTOOTOG CTCACOCCTG TAATCCCAAC ACTTTOOGAG GCCAAGGCAG CTOTOCHDAT GCACCATOCA TOCTCACAGT CCCTTGCCTA TGTGTGGCAG AGTGTCCCAG NTACTAMAA TACGAAATTA GCCAGGTGTG GTGGCACACA TCTGTAGTCC CAG GCGGATCACC TGAGGTCAGG ARTICGAGAT TARTCTGGCC AACATGGTGA AACCCCATNT CACIGCIGIT TITICCICIT IGGICCITCI ATCACIAAAA CICAICICAI TCAGCCITAC TACCICITCI GINAAGCIIT CCCIGGINIC AGGANICAAA AITNAICAGG GAICITITICA GICIGCICIT GIGINGCICA GUAGACAATI CCAGCACAGA CACIACAGIT AACGCIGAAC CAACTACCAT TAGCACTATG TTAGGAGCTG CAAGGCCCCA AAGTAGAAGA TGTGCATAAT AGCATAACTA ATTATTTOTT TICCICACTA CATTOTACAT GIGGGAATTA CAGATAAACG CCAGATOTOT GECCECACEE CATOTECATT TACATOTECT TEAATGEECA CETCAAAAGG AAGIGAGAGA AGCATGAAAA ATGAGCAGGG GCCTGGATCA GTGGGGTGTA TTCAGAGCAC AGCTGGGCCT TGAAGGATGG ATGAAATTTG GATAGAGAAT GAGGAAGACA GAGGGCCTCC TGCAGCTGCA AGTAATAGCA TGAACAGTCA GAAAAATACC TTATGAGGGG GCAGGGCTGA CAMAGACTCA GAGAACTAGA GTTTAAGCTG AGGCAGAGTG CCGCCACCCT GGCATGCCCC TGCAATTAGG CAGATAAAGA CATCAGTCCC AGTAAATGAA TCCATAGACT CATCTAGCAC AAGTATAATC GGCTAACTCC TAAATCCCAA TGAATAGTCC TAGGCTGGAC AGCAATGGGC ICACAACIGA GCAAGACATI CATAIGAICA TITAAGGAAG TOTTICCCTT AIGIOTIAGC ACAAACAGAT CACCAGCCAG CTTACACAGG CATTAACTCT CCTCAATGAG GAAGAATCAT ICTCCAGAAG GAACITIOGGA GATGATIOGTO CAGAITGATIGA AACTIOOGTTC ATCCCAGITIC CIOGOGRACT GGGTATAAIC CAACCAICAA AATAGAAGAC CITOCAAGAA GCAGAGTCAT 3120 3173 3060 3000 2940 2880 2820 2760 2700 2640 2580 2460 2400 2340 2280 2520 2220 2160 2100 2040

ACTECATETE TOCECOOCAG CICTOTOCAG ACATEGCAAT GOGETTEEGT ACTEACEACA

1020 960 900

840

780

720 660

CTGAGGACCA AGAGCTGACT TACACGTGTA CAGCCCAGAA CCCTGTCAGC AACAATTCTG 600

5

TGACATACAA TIGGAGICCC CIGGGAGAAG AGGGIAAIGI CCIICAAAIC TICCAGACIC

CTOTGAACAG CACCTGTAAT GTCACACTGA CATOCTCTGT AGAGAAAGAA GAAAAGAATG

540 480 420 360 300 240

ACAACCIGCA AAICIAICGI CGGCIIGGGA AACCAAAAAI TACACAGAGI TIAAIGGCAI

5

GGATACATIGC CITAGGICCG AACTACAATC TOGTCATTAG CGATCTGAGG ATGGAAGACG CAGGAGACTA CAAAGCAGAC ATAAATACAC AGGCTGATCC CTACACCACC ACCAAGCGCT

CACCAGGAGA CTCAGAAAACA GCACCCGTAG TTACTGTGAC CCACAGAAAT TATTATGAAC

AAGAACCACG GCAAGTTAAA ATCATTOCTT GGACTTCTAA AACATCTOTT GCTTATOTAA

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	405		406	. .	
			ACCTIGANGE CENGENGENE AACEACEATA GEGGGCETEA GGGTEACAGA AAGEAAAGGG	099	
	CCACATTGGA GACTCTGCAG ATCATTAAGC CCTTAGANGT GTGCTGGGTG ACCAAGAACC	300	CACTCAGAAT CATGGCACCT AAGTCTGGAC ACTGCCATCA GGGTTGCATT GGCTGTCGCT	GCT 720	
5		360	5 orgendama engreantiti ogdaenoens tocencence tengrongs aggagaagaa	780	
	TGAGAAAAT CAGCAGCATT GCCAACTCTT TCCTCTACAT GCAGAAAACT CTGGGGCAAT	420	ANGGIAGCAG GGCCCCAAGC AGTGACTTCT GACCAACAGA GTGTGGGGAG AAGGGATGTG	GTG 840	
			_	CCA : 900	•
9	ATGACAACTA TGATCAGCTG GAGGTCCAGG CTGCTGCCAT TAAATCCCTG GGAGAGCTCG	540	 товсьмаят асмтевакаю сиссетсямо иссеттимым озсимлессе самозсивых 	096 CAA	
	ACCIPITICE ACCEPCANT ANTARANC ATGANGANT CTCCTCACCT TCATGACAAG	009	GGNOGCTGGG TCCCTGAATC ACCGACTGGA GGAGAGTTAC CTACAAGACC CTTCATCCAG		
15	GAACCTETAT AGTGATCCAG GGATGAACAC CCCCTGTGCG GTTTACTGTG GGAGACAGCC	660	15 выскитела аспесавтся татаравать аботепваль тослетваят иллаеслеть		
	CACETTRANG GODANGGAGA TOGGGAAGGC COCTTGCAGC TGNAAGTCCC ACTGGCTGGC	720	GCATTIGGGG GCIGITYAIT ATACAGTGC AAAGAGTTCC TITATCCTCC CCAAGGATGG		
	CICAGGETOT CITATICGGE ITGAAAATAG CCAAAAAGTE TACTGTGGTA ITTGIAATAA	780		-	
70	· ACTICIATICTIS CTGAAAGGC CTGCAGGCA TOCTGGGAGT AAAGGCTGC CTTCCCATCT	840	20 TOTATIOGICS CTOTCTICTA TOSCAGAAGS TITITIGGGGAA TAAATAGGGG CANAMENTER		
	AATTTATTGT GAAGTCATAT AGTCCATGTC TGTGATGTGA	006	TCACTHUAAA AABABABAA AABAACTCA		
75	ACACATTGTA CTGAGTGGTT TTTCTGAATA AATTCCATAT TTTACCTAAA AAAAAAAA	960	25	257	
3	AAAAACTCGA GGGGGGCCC GTACCCAATT T	1992		-	
			(2) INFORMATION FOR SEQ ID NO: 177:		
30	(2) INFORMATION FOR SEQ ID NO: 176:	33	30 (i) SEQUENCE CHARACTERISTICS: (B) TENCTH: 2390 base pairs (B) TYPE: nucleic acid		
35		χ. Σ.	000		
3	(B) TYPE: nucleic acid (C) STRANDENESS: double		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 177:		
	(D) TOPOLOGY: Linear		TGGGGCCCCT TITGGATGCT CTGGGTGTTT TTGCCAAGAG TTACAGGATG TCAAGTGTGG	99	
4	(x1) Sequence description: Seq id no: 176;	40	O GENGETCAGE ACCETTOCTO TOGACCAGTS AAGCTOTTE CAGACCAGGT GETTECAGAE	AC 120	
	ACAGGCCTCT TOGGAGCTG AGCCGGCTC TCCTCACTCA CCTCAACCCC CAGGCGCCC	09	ATTTCCAGGC TCCAGGGAG AGGCTGGGAG CCCCACAGA AAGCACAGA AAATGCAAAA	AA 180	
	CTCCACAGGG CCCCTCTCCT GCCTGGACGG CTCTGCTGGT CTCCCCGTCC CCTGGAGAGG	120	AAAAAACAGF CITITITITI TITITIGCITT TIATIAIGAA AACAAAACAA AIGCCCCAGG	240	
45	AACAAGGCCA TGGGTCGGCC CCTGCTGCTG CCCCTRCTGC YCCTGCTGCM GCCGCCAGCA	180			
	ITTCTGCAGC CTROTGGCTC CACAGGATCT GGTCCAAGCT ACCTITATGG GGTCACTCAA	240	TOTTGAGGOC AGCATTGCAA TAAACAAGCT AAACTACTTA CATTGGACTC ATTTTCAGTA		
20	CCAAAACACC TETCAACCTC CATGGGTGGC TETGTGGAAA TECCETTETE CTTCTATTAC	300 50			
	CCCTOSGAGT TAGCCAYAGY TCCCRACOTG AGAATATCCT GGAGAGGGGG CCACTTCCAC	360	AACTTGGATG CACATCATAC AGAAAAGTAA CATTTTAAAT ATAAAAAAAA AAAACTTGCT		
	GOOCHOTICT TETACAGCAC AAGGCCGCCT TCCATTCACA AGGATTATGT GAACCGGCTC	420			
55	TITICTIGNACT GENCHGAGGG TCAGGAGAGC GGCTTCCTCA GGATCTCGAA CCTGCGGAAG	480 55			
	GAGGACCAGT CTGTGTATTT CTGCCGAGTC GAGCTGGACA CCCGGAGATC AGGGAGGCAG	540	AAAACAAACA CACAAAAAA TOTOTOTTAC AGTITIOTAAG CAAGATGACA CTGCCGAAG		
9	CASTIGGAST CCATCAAGGG GACCAAACTC ACCATCACCC AGGTISTCAC AACCACCACC	09		•	

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50 45 8 33 30 25 20 15 5 S CCTCCACCCC TOGGGAGGGC AGACAGGCTC GGGARGGCCT GGCCAGGCCA CTGGAGGCTG ACAAAGTITA TATTATATAA CIGGOGTICC CIAAATIGAT TICTITIAAA ACAGICITAA GCAGGGAGCA GGCATGTCCA CCCGCAAGCC TGGGAGGCTA ACTCTGGCAT TCCTGGCCGG GCGTGAGCAG GGAGCACCGT GCGAGTCTCC GGGAGGGAAAT CCTCCTGGGG CCCAGAGACT ACCINCAICT GIANGOGICA GIOGACICIG ANICANTITI AIGGIIGITI TAAAAICACC ACTIVATIVA ANGITTANCA ANCIGGOTGA ANACTOROCA AGTOTORGAC TORCONGCA CGAGGGTAAC TITAAAAAAT GGAAACTITC AAATCCATIT ATATTITIAT TATAAACAAA CIGIOCIOGO AGCACGITAC CAACCAGCCI GCGIGAAGAC CIGICAACIG ICGIGIGIGA GOCCOCTECE COGACOCCTE ACACAGOCAG CACCTCACTO CECTOTOGET GGAGOGGCAT CAGAGAACCA GCCGAGAAGG AAAGGCCCCA CGATGCTCCC TGTGCGCTGC CCCCACAGCC CAGAAAGCCT TAAAAAAGTG ACAGCACCAA TGCAGCTGCT CAGTGTACCC NCCGTGGGCT TGAAAGCTGT AGCCTCTGGG AAAACAAAAC CAAAACATCA CCTTCTATTA AACTCTGTAT ATTATTATTT AGCCGCCATG CTCATTGGTG GGCCAGTTTG GGACATCCCC GTACTCAAAG ACCATATGGC CCCACGETGA GETCCGGGAG AATGCCTGGT TTCAGTCATT TCCGGACTAA CTGTGACAAC GECCACCATA CAGGACAGAC CACACCACAG CTCCATACCC AGCGTCTGCC TGGAGGCTCC GTGTATTAGG ATACTAATGA TAGTCCCTAT ATCCATCCAG AAATGCTGGC AGAAAGCACT TTTAAAAAAT GATAATITAC CAGCATCICC ICATCAGAGT ICCCICICCA GTAAGGGIAI ATTOCTTAAA TIOGGITIAA ATAGIOCATT AAAGATOTGT TIAGAAAATA COTTIGAAAA TOCAMOUNDO OCCCCCAMO CCCAMOGCACO CCCGGCTTAG GOTOTACIOTA TCACCCAMOC GTCAGGGTCA GTGGCTTCTT TCTAGATGAA AGGAGCAGAG GCGAGCCGAC GCCACCGTCA GCGCTGAGGA AGTCATTAAT CCTTCGAAAAC TCTGAAAAGA AACCAGTGTT GAAGTCTGGA CCGCAGCCTT CTAATACAGA AGAAACGGAC GTGACTGTCA CCCTCAGCCC GCCAGCAAGG NGAGACCAGA AGTGAATACA AAAGAACTAA ACAAAATAAA AAATTAGAAT GTGCTGTAGC ITTACAATAG AAAGITAAAA AICAAGACIT AGAITTACIA TACAITTITIT CICICAGATI NAMAMAGCAT GTTAGAAGCT GCCCTACAGG TCTCAGCAGT GGGACAATCT AATTGAATCA ATTICAGAGTG TICTCAAATC CAATICCGAC ACACGACTIG ICACTACTCC ICTCCCCTTG 2280 2100 2040 1980 1920 1860 1800 1740 1680 1620 1560 2290 2220 1500 1440 1380 1320 1260 1200 1080 1020 960 90 840 780

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20 2 5 S GGCACGAGCC AUGCCUGGCC TCUCCTUGAT TCTTACAGUC ACUTUGUIGG CUGUTUCUGA IGITICCAAG AIGCITCIGA AGATIGCCIA AAAATAGCCG GITTCCACCC CCGIQAATGC AGTAGAGAGG TAAAAGGCCA CCATCTCCTT GACCTCTGGG GAACTCATCC ACAAAGAAGA THOSTOCAGO AATOTAGTAG OCATACACOT GOTTOCGTOG ATCTOGGCCC TCCTGATOTO ATCAGTAACA ACCAGGAGAG AAGCTGCTGG AACTGACCTC TOOGAACTCC CTGGGATGGT TGTGGAATAG TGTCTGTCCA TGCCTCTCCT CATGGGCTAC CACCTCTGCC ACCGTGGTTA CICAGCAGCI ACCIGCATIG IGGCCAAAGG AIGACCIATI CCIICICAGG AGGGCAAAAA SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 549 base pairs 178:

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SEQUENCE CHARACTERISTICS:

SEQUENCE DESCRIPTION: SEQ ID NO: 179:

(D) TOPOLOGY: linear (B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1509 base pairs

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INFORMATION FOR SEQ ID NO: 179:

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CCCAGAATTT GCAGTAGCTC AAATTAAGTT TCTAGCTATT AAAAAGAAA GAATGCTCCT TCACCAGGAC CAGAGAACTG ATTTACAGAA GTGACATGAA

549 540 480 420 360 300 240 180 120

aaaaaaaaa AACATTCCAT ATCCATTCTA

CAACAGCCAG AACGGGGAGC GCTGTGTGGC CGCCCTGGCT CGTGTCGAGC CGCCATCCAG ATCTGCCGGA TCATGCGGCC AGATGATGCC AACGTGGCCG GCAATGTCC COGGGGGACC ATCCTGAAGA TGATCGAGGA GGCAGGCGCC ATCATCAGCA CCCGGCATTO GOODGCOCA TOOGCOGCOG CAGOODCOCAG CATOTOGGGC CCAGACOTOG AGACGCOOTO GGCACGAGGG CTCATTCATT CCGCGCCGGG CCTGCCAGAC ACCTGCGCCC TTCTGCAGCC GCACCGACTI 240 180 300 120 8

3

GCGGTATGAA GCCCAGAAGC TOGAGCOCAT GGAGACCAAG TOGAGGAACG GOGACATCOT GOTOCTICGAG GIGCCTICCTG TIGTGTATTIC CCGGCANGAG CAGGAGGAGG AGGGCCGGAA AMBOTTEMOC AMENAGENCA COUTTETETA TOTOCCOCTO TOGOTTEMAGA ATOTOGRACA CAAGCACTCT GIGGROGICC AGGICARCGI GAIGICCGAA AACAICCICA CAGGIGCCAA 540 600 480 420

COTOTICICO ATOTIOCATOS GTUAGGIOGO GCATOTICAGO GCGGAGATICA COTACACOTO

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INFORMATION FOR (SEQ ID NO: 178:

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	409			410	
				TITICAGISC COCACARISA GICAGAAGAT GAAGTGGCTG TICACATGGA ATTTGCTAAG	480
	CCAGCCAGTC CTCAACCCAG AGOGGAACAC TOTCAGCTAC AGOCAGTCCA GCTTGATCCA	099		ANTATATATO AACTICCATAA AAAAGTTTCT. CCAAATGAGC TCATCCTGGG CIGGTACGCT	540
v	CCTGGTGGGG CCTTCAGACT GCACCCTGCA CGGCTTTGTG CACGGAGGTG TGACCATGAA	720	2	ACOGGICATO ACATCACAGA GCACTICTOTO CTGNATCCAT GAGTACTACA GCCGAGAGGC	009
	GCTCATEGAT GAGGTCCCCG GGATCGTCSC TGCACGCCAC TGCAAGACCA ACATCGTCAC	780		CCCCAACCC ATCCACCTCA CTGTGGACAC AAGTCTGCAG AAGGGCGGCA TGAGCATCAA	099
			9	ACCTACOTC ACACTITA TOGRAPICC TGGGAGACC ATGGGAGTGA TGTTCACCC	720
2	CTCGGBACGC ATGACCTTCA CGAGCAATAA GTCCAITGGAG ATCGAGGTGT TGGTGGACGC	006	2	TETGACAGTS ANTREGGET ACTAGGACAC TGAAGGCATC GGAGTTGACC TGATCATGAA	780
	CGACCETOTT OTGGALAGET CTCAGAMOCG CTACCGGGCC GCCAGTGCCT TCTTCACCTA	096		GACCTICCTIT ACCCCCAACA GAGTGATTGG ACTCTCCAAGT GACTTGCAGC AAGTAGGAGG	840
15	COTOTCOCTG ACCAGGAAG CCAGGTCGCT OCCTGTGCCC CAGCTGGTGC CCCAGACCGA	1020	15	GECATCAGCT CECATCCAGG ATGCCCTGAG TACAGTGTTG CAATATIGCAG AGGATGTACT	006
	ggacgagaag aagcocttto aggaaggcaa aggocggtac ctocagatga aggogaaggr	1080	Ū	OTCTORANA GISTCAGCTG ACANTACTOT GGGCCGCTTC CTGATGAGCC TGGTTAACCA	096
	ACHORGOCIAC GOGGASCITIC AGCOCINADAC TOCOTICOTICO TIGOCACTIGOTI GOCTICOMATIA	1140	` ?	AGTACCGAAA ATAGTTCCCG ATGACTTTGA GACCATGCTC AACAGCAACA TCAATGACCT	1020
20	OCCATGOCAA COSSOCCAOT OTCCAGTCAC TINGAAGTTC COCCCTTGGC CAAAAACCCA	1200		TITIGATIOSTS ACCTACCTES CCAACCTCAC ACAGTCACAG ATTGCACTCA ATGAAAACT	1080
	ATTCACATTG AGAGCTGGTG TTGTCTGAAG TTTTCGTATC ACAGTGTTAA CCTGTACTCT	1260		ISTADACCTG TGAATGGACC CCAAGCAGTA CACTTGCTGG TCTNGSTATT AACCCCAGGA	1140
25	CICCIGGAAA CCTACACACC AAAGCTITAT TTATATCATT CCAGTATCAA TGCTACACAG	1320	25	CICAGAMOTO AACGAGAAAT GOOTITITITG TGOTCTIGAG TCACACTGAG ATAGTCAGTT	1200
	TOTTISTICCO AGGECGGGA GCOTTIGGC AGAAACCCTC GGGAATGCTT CCGAGCACCC	1380	G	GIGIGIGACE CTANTARAGG GAGGCTACCE TITIGIFAANTE AAAAAAAAA AAABAAACCH	1260
	TOTAGGOTAT GOGALGALCC CAGCACCACT AATAAAGCTG CTGCTTGGCT GGAAAAAAA	1440		SGRGGGGGG CCCGGTCCCA TTSSCCCTTT NGPAARTYCT NTHACAARTC	316
30	ANANDARAA AABAHRAAA AABABARAA AABABAAA ABABABARA ABARABAR	1500	30	SOLITO	9177
	Agaaaaan	1509			
36		6	35 ((2) INFORMATION FOR SEQ ID NO: 181:	
Ç	(2) INFORMATION FOR SEQ ID NO: 180:				
Ş	(i) SEQUENCE CHARACTERISTICS:	4	40	(c) Tyrk: nucleic ecid (c) STRANDENESS: double	
\$	(A) LENGTH: 1316 base pairs (B) TYPE: nucleic acid (C) STRANDERNESS: double		•		
	(D) TOPOLOGY: linear	•		GOCATOWICA GACATGACTT CTATTGCCAG GCTGGTCAAG TGGCAGGGTC ATGAGGGAGA	09
45	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 180:	44	ნ მ	CATCGATAAG GGTGCTCCTT ATGCTCCCTG CTCTGGAATC CACCAGGGG CTATCTGGGT	120
	ACCITITATICA TIAGGAAAGAT GOCCACACOG GCGGTACCAG TAAGTGCTCC TCCGGCCAGG	09	E	TIATGGGGCT GGGGACTAGA ATTGGATGCT TCAAAACCAT CACCTGTTGG CCAACAAGTT	180
90	CCAACCCCAG TECESGOGG GOCCCAAGC TCAGTTCCAG CGCCAACGCC AGCACCGGCT	120 5	50 n	TGACCCAAAG GTAGATGATA ATGCTCTTCA GTGCTTAGAA GAATACCTAC GTTATAAGGG	240
	GOGGCTCOGG TTCCCGCTGC GGCTCCAGCC TGCATCCTCA GACCCTGCGG CAGCAGGGGC	180	8	CONTICTATE OCGACCIGAA CITTGAAGAC CACAMIATIG AAGAGGGGTE GCTTACCYGT	300
	TOCAACTEGO GCTCCTGGCC AGACCCCGGC CTCAGCGCAA NTOCAGCGCA GACCCCAGCG	240	•	TOOGOCCAA GAGCATGTT ACCAAACATG GYYCARGAAM YTTOGYKGGG AMCARIGGGKG	360
55	COCRETICIOS CITAGOCIOS ICITACARAS COCITACOCOS ACABOCADOS ASTONABETA	300	S B	GENGOGAREM CHRICGYTTG SCAANTTCSK KGOCHNCCYT TTAGGGTAAR RRGGGCKGTW	420
	CACCCAGTICA TITTIGGCCTC CATTGTGGAC AGCTAGGAG GACGCAAGGA GGGTGCTGCC	360	A	ATTAGATICT GOSTAAASTA GGAICTITTIS COCTIGGAAA TITIGCITGCC GGGIGAAATS	480
. 09	CONCITATIOG GOACCOTOTT GOGNACTOTO GACANACACT CAGTGGAGGT CACCAATTGC	420 66	وں بو	TOCTTOTICC TICHORACCE CIPACCEING TAGTICCTICS ACTIVACITIC TCACIANOTIC	540

8 Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1596 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double

55 (2) INFORMATION FOR SEQ ID NO: 184:

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TAAAAAAAA AAAAAAAAAA CTCGA

1405

GATTOGTGTA AATCITTITA TAAATACATA AATAAAAGNA AAATATGCAT TITITCTTTTC

TCATATATIT AACTITGCAA AAAGATITAC TITGTACAIG TTACAGGCTT

1320 1260 1200 1140 1080 1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 69

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GGGAAACTAA TIGIATITAG TGACAAAAAT AAAAAGTITI TITITIATAA

TICAGICIGO

CTOTACAGAC CATCTOTATG TTAGGTGACA TIGATTATGG GTTATAATCA TITATCTGAG TITAGTGGTC CTAATATATA TGTAGAGAAA GATGGTGGGG ATTTCCTTTA AATGGTAAGA GTTTCTAAAA CAGACAATAA TTTAACAAGC

TITIGGATIT

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AAGATTATTT TATCACTACA AGCTTTTAAC TTTTTAAGTT ATTGTACAAG TATTCTACCT TITICIAGAC TITOGGATCTO CAAGAAGGCC AATTOCCTAA AATTICIGAG AACAGIGCAC CACCACGACC AACCATATGO CAAATGAACC AAGCCCAGTT GITGCAGTGA TIGGITGICT 30

TATATOSCIT TECTTCACCT CICCOTCATG ATTOTICTOT TGACTTACAC ACCAGAAATG

AGTATTCGCC TOGGAATTTT TCTCCGAAGA TACCCCATAG COCGAGTTTT TGTAATTATA ACTAATCIGG CAGGAATOTA COGAAAAOTT COCAAAGCIG CIAGIICAAT IGAICAGIIT ATTGACAATO GTGAAGGCAC TOGICTOCGA AATGTTCCTG TICTTTTTAA TGACACAGAA CAGCAGATGA ACTOCOCCTO TOGAAGTAGT AGTAATOGGT CTTOGATTAA TATGTCTOGA ATGCTGGAGA GTCTCAGCAC AGAAAAGAAC TCCCTGGTCT TTCAACTGGA GCGCCTCGAA CAGTOTOAGT TAGAAAATOG ACTOCATOAG CTAACAGAGA CTOTOATOCA GAAACAGACO GACGAAGAAA TTCAAAAACT CAGGAATCAG CTTACCAATA AAACTTTAAG CAATAGCAGT TATATAGAAG AAGATCTTTA TCGAACAAAG AACACATTGC AAAGCAGAAT TAAAGATCGA CAGAAAGCAT CCAAACAAGA ACTAGAGACA GAACTOGAGC GACTGAAGCA GGAGTTCCAC GTTAATGAAG CAGAATCAGC AAGAGAACAG TTACAGGWTC TGCATGACCA AATAGCTOGO AAGCTGATGG GCCAGATACA TCAGCTCAGA TCCGAATTAC AGGATATGGA GGCACAGCAA

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AAATTGATTA ACAGCTTGAA AGAAGGCTCT GOTTTTGAAG GCCTAGATAG CAGCACTGCC agtagcangg agcnogaaga actnogocan gagaaagaga ngcagaggga ggaaanacag

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 183

INFORMATION FOR SEQ ID NO: 183: SEQUENCE CHARACTERISTICS: (A) LENGTH: 1405 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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2 8 35 50 ဗ 25 TOGITICITY CCICCATACC CCITCIGCAT TICAGIGITY TIGITYAGIT TICCIGGITT CCACTAATAG AATTCAGCTT TTAGCATGGG CTGTTTCATA CTGTTCTGAT GAAACTGATT CACCACACCT ACAGGGACCA CGTGGTGGGC TGTGGACTAG CGGCCAAGCT CCCTGCAGGC CCTGCACCAC CITGAAAGAC ATTICTAATA TGGTTIGICA GGCAAAGIGG TTAGIAGICA ANANANACT C TOCCCCTOGA CTOGGCTCTG TOAGAGTOGC CTTCTGCACT GTGCACAGTA GGTGTGAACA GATCYTGGGA AAGANATATC TIGCCAGGAA AAATGATAGN CCTIGACAAT GIIGAATGAT ACTOGICTIC AAATOTOTAC ATOTOTOCCA GOGAGCAAAT GCCTTCTTGT TICTGAAATT ITTIGTGGCCT GAGGTAGAAG TCCTCAGAAA TCAGCAGACT TCACTGATAA AATGCTGACT ACATGTOGGC TOGAGATCTA TICATITICGT TITIGGCITGA ATTITICTGRA TGGTTTACTT TECGTGGECA GAACCECTEC AGGTEAGAGG CAGAAGAGAA GCCTCATGGG TCACAGCAGC GGICTITIAG ACTGITCITI TITICCCANCT TCTCACCICC TGCCCCCCCCT TCAGGGIACT GGCACAGATA ACTATOTACA TOTATTCCTT AAATGITTIT TIAAGITITA TAITCTIGGC 791 780 720 660 600 540 480 420 360 300 240 180 120 60

SEQUENCE DESCRIPTION: SEQ ID NO: 182: TOPOLOGY: linear

20 15 5 3 INFORMATION FOR SEQ ID NO: 182: (1) SEQUENCE CHARACTERISTICS: (C) (E) (A) LENGTH: 791 base pairs TYPE: nucleic acid STRANDEDNESS: double

agaatgagaa ctoctotgat agogagagto aaggagggat atotogtaga gcacttgatt 777 720 660 600

AGGITITGGTA GCGTGGAGGA GAACIFITGAT GGAAAGAGAA CCTTCCCTIC IGTACIGITA TCAGTTGAAT GCCTGCTGGT AGCTTTTCCA TTCTGTGGAG CTGCCGTTCC TAATAATTCC

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(2) INFORMATION FOR SEQ ID NO: 185:

. 8 120 180

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 184;

S	STCATISCAST GOBOCOGRIGA ACTISTISCITOT TITCAGOCOGA COCTIAGGOGO COCGAAGOGA	09
	AACTIGGGAGG GGAAGGTGAC GGGGAACGGA GCATTTCAGA TCTGCTCGGT AGACCTGGTG	120
2	CACCACCACC ATOTTGGCTG CAAGGCTGGT GTGTCTCGGG ACACTACCTT CTAGGGTTTT	180
2	CCACCCAGCT TTCACCAAGG CCTCCCCTGT TGTCAAGAAT TCCATCACGA AGAATCAATG	240
	GCTGTTAACA CCTAGCAGGG AATATGCCAC CAAAACAAGA ATTGGGATCC GGCGTGGGAG	300
13	AACTGGCCAA GAACTCAAAG AGGCAGCATT GGAACCATCG ATGGAAAAA TATTTAAAAT	. 360
	TGATCAGATG GGAAGATOGT TTGTTGCTGG AGGGCTGCT GTTGGTCTTG GAGCATTGTG	420
2	CTACTATIGGC TTGGGACTGT CTAATGAGAT TGGAGCTATT GAAAAGGCTG TAATTTGGCC	480
2	TCAGTATISTS AAGGATAGAA TTCATTCCAS CTATATISTAS TTAGCAGGGA GTATTGGTTT	540
	AACAGCTITG ICTGCCAIAG CAAICAGAG AACGCCTGTT CTCATGAACT TCATGATGAG	009
ς:	AGCCTCTTGG GTGACAATTG GTGTGACCTT TGCAGCCATG GTTGGAGCTG GAATGCTGGT	099
	ACGATCAATA CCATATGACC AGAGCCCAGG CCCAAAGCAT CTTGCTTGGT TGCTACATTC	720
ç	TOSTOTISATO GOTGCAGTOS TOSCTOCTOT GACANTATTA GOGGSTOCTO TICTCATCAG	780
•	AGCTGCATEG TACACAGCTG GCATTGTGG AGGCTTCTCC ACTGTGGCCA TGTGTGCCC	840
	CAGIGAAAAG TITICIGAACA TGGGTGCACC CCTGGGAGTG GGCCTGGGTC TCGTCTTTGT	006
S	OTICITEATTO GGATCTATGT TICITICACC TACCACGTG GCTGGTGCCA CTCTTTACTC	096
	AGTGGCAATG TAGGGGAT TAGTTCTTTT CAGCATGTTC CTTCTGTATG ATACCCAGAA	1020
	AGTAATCAAG CGTGCAGAAG TATCACCAAT GTATGGAGTT CAAAAATATG ATCCCATTAA	1080
,	CICGATOCTG AGTATCTACA TOCATACATT AAATATATTT ATGGGAGTTG CAACTATGCT	1140
	GOCAACTUGA GOCAACAGAA AGAAATGAAG TGACTCAGCT TCTGOCTTCT CTGCTACATC	1200
S	AAATRICITG TITAATGGGG CAGATATGCA ITAAATAGIT TGTACAAGCA GCITTCOTTG	1260
	angtitagna gataagnaag atgtcatcat atttaaatgt tecggtaatg teatgectca	1320
c	GGICTGCCTT TITTICTGGA GAATAAATGC AGTAATCCTC TCCCAAATAA GCACACAT	1380
,	TITICAATICT CATGITICAG TGATITIAAA ATGITITIGGT GAATGIGAAA ACIAAAGITT	1440
	GIGTCARGAG AATGIAAGIC TITTITCIAC ITTIAAAKITI AGTAGGITCA CIGAGIAACT	1500
S	AAAAITTIAGC AAACCIGIGI ITGCAFATTT ITTIKGGAGIG CACAAFTANIG TAATTARAGC	1560
	ATTCCAGTAA NAGTGTNTTT AAAGTTGNTC TATAIN	1596

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240 300 360 450 480 540 909 999 720 780 1140 1200 1260 1380 840 900 960 1080 1320 1440 1500 GOSCAGAGOC COYACGAGCA GGACGACGAC GACAAGGGOG ACTCCAAGGA AACGCGGCTG ACCCTGATGG AGGAAGTGCT CCTGCTGGGC CTCAAGGACC GCGARGGTTA CACATCATTT TGGAATGACT GTATATCATC TGGATTACGT GGCTGTATGT TAATTGAATT AGCATTGAGA GGAAGGTTAC AACTAGAGGC TTCTGGAATG AGACGTAAAA GTCTATTAAC AAGAAAGGTA TGGAATCCAT TAAAATTGCA TTATCAGTTA AGAAATGTAC GGGAACGATT AGCTAAAAAC ATCTGTAAGT CAGATGCTCC AACAGGGGAT GTTCTTCTTG ATGAAGCTCT GAAGCATGTT ACACATCCCC TCACCAATAA CAACATTAAG CAGCGCCTCA TCAAGAAAGT ACAGGAAGCC CITICITICACA AATGGGTGAA TGACCCTCAC CGCATGGACA GGGGCTTGCT GGCCCTCATT AAGGAAACTC AGCCTCCAGA AACGGTCCAG AACTGGATTG AATTACTTAG TGGTGAGACA INIGATITIOG CTACCAAGAG AGTGCGGCAG CTTCTCGACT TAGACCCTGA AGTGGAATGT CTGAAGGCCA ACACCAATGA GGTTCTGTGG GCGGTGGTGG CGCGTTCAC CAAGTAACTC IGCTCGGGGF GAACCAITICT CCTITICTICTC AAGTAAACCA GTAGTITITIC TICTGITIGAC TICTIGGITIT CIGIAAITIG TACTITICCCA CACTATAATT GGCTTCTGIT TTACAAAATG TACCTIGACTO ATGOSTOGGA COTOCTIGGAG AATGOTTTTIG CTCCTCTTCT GGACGAGCAG STOGGTOSCT TITICITITY TOTACGIOTA CAGALTICTO CIOGTACGAG AGGCCTTCCT CITICIOTIT TIABABABG ITTIBCIGC ATAITIGGCAT ICCATICCCI GITCCCAICC TCACTGTTAC CTGTTTTGGG TITTCTGGTCT ACTITTGACTT TCAAAGTACC TCCAGCCTCC TCATAGGGAC AGCTTTTGGA TGACCTCAGC TTGAGTTTCT CCATATGTGC ATGTACATCT AGCATTICTIGE CTACAGITICA GACAGAAGTIC ACAAAAAGGE CTTCAACTICA CCAAAGGTAA ATATOTOTAT CTATTAGGAC ATTITITACA TAGACTICAG TICAGATGIA TACTTAGGAA AAITAITITI AAAITGAAAC AGCACAGIAA ATACITAATA TAAAANGICC CITGGAITITI GCTTCCCATG TAAATCTATT GTATTATAC ACTIGTTATA ATTTTAACTA TAAAGGTCCA ATTOTITICAC AGAGCCAGIT TGGGATGGGC TGCATTCCAT TTATGCTGTA TATAGTTTGA · ATTATATATA AATTACCCCT TCTTCTGGCC ACCCTGCTC CCATCTTAGT ATTTTGCAAG SEQUENCE DESCRIPTION: SEQ ID NO: 185: (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2293 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear (x <u></u> 2 20 25 30 6 8 35 45 20 55

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ATCTAATCAG TIGIACACCT GGIGCCCCTC GCTIGCITCA ATCAIGGITA TITIGAIGGCA

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aaaaaaaaaa aaa GTTAATAGTG GITTITAATIT TITIATATACA ATTAGGATAA TIAGCATIGI CAGACTATAA ACCITIGCIT TACTCACACA TICTIGOGGA AAAAAATCAA AIGICAGICC TAGCAGAIGI IGCAIGIAAA TCTTGAGTTT TAAATTITAT CITACIGITT TIATAATTIC TATIOGGAAG AGGCTIGIGA CCAGTACCAA TITTIGAGOT GIGCCATTIA TOGIACIOTI TOCCTATGCA TOCCCTITIT AGATITITI ACTATOTITT TATOTGAAGT AATAAATAAA CAATGATOTT GAAAGTOOOY RAAAMAAAA AAAAGTACTA GATGGTGTAT AACTCTAGAG TIGAATTITTA AGGGATTCCC TAATATGTAT TTTAAAGTTT TTOOTAGCAA GTAATGATTA CAACCCAGAG GATTAAGAAT TTTGTAACAG AAAGCTCTAT AAATCGACCT CTTGTCGCTG AAGGAGAGAG AAAAGATGTG TGTCTGATTG GTCCTGGGAT GGGCTTTCAG CTATATOTTO AACAAATTAA ATGTCAAAAT TITTTATTAC CATAGTCCAT ATTITITACIA TENCITTATC ACTITATIST ATCATCACCA TIGGIFTICAT CITITITCIGI CCACAAGIAA AITAAIAICI GCICIGAAAI GICATITAIC GIGITIAGAG AUTITITITIO ITGITOTIAA CATICATIOC

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INFORMATION FOR SEQ ID NO: 186:

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Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1212 base pairs

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45 CCGTGGGTTT GCGACGTTTA GCGACTATTG CGCCTGCGCC ACGCCGGCTG CGAGACTGGG GOCACGAGGC GAGCCGGCGC ACCGTACGCT GGGACGTOTG GTTTCAGCTC GTGCGCCTCC 180 120 60

GTGTAGGTGG GOOGRACIA CIBGIOCOGO GIBAIBOTAG GOGGOOCOCO GOGGIOCAGO CIBTIBOOGO GAGTCACGGA CGGTTCGGGG CCCGAGGTGT CCGCGAAGGT GGCGCACATG

GGACATIGAG AAGGACCAGA TINITGAGAT GGCCIGICIG ATAACIGACT CIGATCICAA GECCCCAGGG GAGAGCATGG CTCAGCCGAT GGTCTGGGTG GACCTGGAGA TGACAGGATT

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CATTITIGGCT GAAGGICCTA ACCIGATTAT AAAACAACCA GATGAGITGC IGGACAGCAI

STCAGATTOG TOTAAGGAGC ATCACOSGAA GTCTOGCCTT ACCAAGGCAG TGAAGGAGAG

TACAATTACA TIGCAGCAGG CAGAGIAIGA ATTICIOICC TIIGIACGAC AGCAGACICC

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ATCCTACTCC ATGGGATGAA GACGTCTGAC ACTATTATCC GGGAGGGCAC CCTGATGGGC CCCCAGAAAA TIGCAGGIGA ACICIAIGGA CCICICAIGC IGGICIICAC ICIGGIIGCI

720 660 S

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480

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 186:

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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CCCACCIOTO CICGACCOTO GACGICIACO TICOGGAGGO CCACATOTIG CCCACIOCOC GCTTCCGGAA GTTGCTTTTG TCCAAACATC CGGGCTTCTC CTTTTTGTGT GOSCOGOGG GCGGCGCCG AAATGGAGCT GGCCCGGAAT GGGGAGGGTT CGAAGAAAAC GCOCOGOGET AGCOCOGOTT TCAGEGAEGG GAGCCCTCAA GOGAEATGGE AACTAEAGCO TCCGGCCGAT

ATCCAGGGG GAGGCTCAGC TGTGATTGAC ATGGAGAACA TGGATGATAC CTCAGGCTCT 240 300 180 120

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GCAGCTGATG CAGCTGCTGC TGAAGAGGAG GATGGAGAGT TCCTGGGCAT GAAGGGCTTT AGCTICGAGG ATAIGGGIGA GCIGCATCAG CGCCIGCGCG AGGAAGAAGT AGACGCTGAT 420 360

TOCAGGOCOT TOAGOTTIGTA COCCAACATO GACATOCTICA GACOCTACTT TGATGTGGAG AMOGGACAGO TGAGOOGGCA GOTGGCAGAT CAGATOTGGC AGGOTGGGAA AAGACAAGOO 540 480

CCTGCTCAGG TOCGAACAGG GCTCCTOGAG TCCATGATCC CTATCAAGAT GGTCAACTTC 600

2280 2160 2100 2040 2220 1980 1920 2293 1860 1800 1740 1680 240 1620 1560 420 360 300

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2 INFORMATION FOR SEQ ID NO: 187:

E SEQUENCE CHARACTERISTICS: (A) LENOTH: 1605 base pairs

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(B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO:

187:

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aaaaaaaaaa an ТИСТТСТИТТ СМАЛТССАЛА АЛЛАМАЛАЛА ЛАЛАЛАМАЛ ЛАЛАЛАМАЛ ЛАЛАЛАМАЛ TATATICCATT TOCTITITANA CCATTITCTTT TOTTTANATA AATAAATAAG TAAATAAAGC

1200

8 TITICCATTAT GACACAGCAG CICCITIGIA AGTACCAGGI CATGICCATC CCITGGIACA

2 CTCCAGATTG ATTACTCAAG CAGACAGCAC ACGAAATACT ATTTTTCTCC TAATATGCTG

1140 1080 1020

TOTOGIOGIT TITITITICIC ACCORGANGS CITOGCAGAG CACCITOGGI TAACITGCAT 960

900

GANGACCOTO ACTIGATICC ACTIVICATO CICCCACIAC AICCITIAICI OGAGOCAACI CATCTTCAAG AAAAAAATAG ATGAAAAGAA GAGGAAAATT ATAGAAAATG OGGAAAATGA

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ACCGAAATAA

ACCCIGCITO

TCATAGGGCA CTTGATGACA TTAGTGAAAG CATCAAAGAG CTTCAGTTTT AGAACTOTIC AGACGCTOOT ATCCAGAAGA ATATGAATTT GCACCAAAGA ATACATOCCC CAGTICATGA AACATCTICA TTATAGAATA ATTGATGIGA GCACTGITAA ICCAGGOCTC TGTCCACTTG CAGGAAATTC AGTTCATGAA GATAAGAAGT TTCTTGACAA 720 660 780 8

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780	840	006	096	1020	1080	1140	1200	1260	1320	1380	1440	1500	1560	1605					09	120	180	240	300	360	420	480
ACAGCAITG GCACCTGCTT CGGCTACTGG CTGGGAGTCT CATCCTTCAT TTACTTCCTT	GOCTACCTOT GCAACGCCCA GATCACCATG CTGCAGATOT TGGCACTGCT GGGCTATGCC	CTCTTTGGGC ATTGCATTGT CCTGTTCATC ACCTATAATA TCCACCTCCA CCCCCTCTTC	TACCICTICT GOCTOTTGGT GOSTGGACTG TCCACACTGC GCATGGTAGC AGTGTTGGTG	TOTOGRACIOS TOGGICCICAE ACAGOSSOTIS CTOCITOTISTO GCACOCTIOSC TOCOCTIACAE	ATGCTCTTCC TGCTCTATCT GCATTTTGCC TACCACAAAG TGGTAGAGGG GATCCTGGAC	АСАСТИСАНО ССССЕЛАСАТ СОСВССЕЛТС САСАВЛЕТСЕ ССАВЛЕНСАТ СОСТВСЕЛТБ	CTCCCTGCTG CTCGGCTTCC CACACGGTC CTCAACGCCA CAGCCAAAGC TGTTGCGGTG	ACCTIGART CACACTGAC CCACCTGAAA TTCTTGGCCA GTCCTCTTTC CCGCAGCTGC	AGAGAGGAGG AAGACTAITIA AAGGACAGTC CTGATGACAT GITTOGTAGA 1000GT1110C	AGCTGCCACT GAGCTGTAGC TGCGTAAGTA CCTCCTTGAT GCNTGTGGGC ACTTCTGAAA	GOCACAAGG CAAGAACTC TGGCCAGGAC TGCAAGGCTC TGCAGCCAAT GCAGAAAATG	GOTCAGCTCC TITGAGAACC CCTCCCCACC TACCCCTTCC ITCCTCTTTA ICTCTCCCAC	ATTGTCTTGC TAAATATAGA CTTGGTAATT AAAATGTTGA TTGAAGTCTG GAAAAAAAA	adaaraaa aaaaaaaaa aaaaaaaaa aaaaaaaa cacaag	(2) INFORMATION FOR SEQ 1D NO: 188:	(1) SEQUENCE CHARACTERISTICS:	(A) LENGTH: 1516 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 188:	ATTUGGCATG AGGGGGTCAC GTGGTGGCTG GGCCGGGGAA ATGGCCGCTT CAGGAGAGA	COGRACTICA GOOGGOGAG GCAGCACCA GGAAGCATIT AIGACCITCT ACAGIGAGT	GAAACAAATA GAGAAGAG ACTCGGTTCT AACTTCGAAA AATCAGATTG AAAGACTGAC	COSTOCIOST TOCTOTIACT TOARTITGAA COCAITIGAG STIVITOAGA TAGATOCIGA	AGTTACAGAT GAAGAAATAA AAAAGAGGTT TCGGCAGTTA TCCATCTTGG TGCATCCTGA	CAAAAATCAA GATGATGCTG ACAGAGCACA AAAGGCTTTT GAAGCTGTGG ACAAAGCTTA	CAACTIGCIA CIGGATCAGG AGCAAAAGAA GAGGGCCCTG GAIGIAAITIC AGGCAGGAAA	AGAATACGTG GAACACACTG TGAAAGAGCG AAAAAACAA TTAAAGAAGG AAGGAAAACC
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540	9009	99	720	780	840	900	960	1020	1080	1140	1200	1260	1320	1380	1440	1500	1516	
TACAATTOTA GAGGAGGATG ATCCTGAGCT GTTCAAACAA GCTGTATATA AACAGACAAT	GANACTETT GCAGAGCTGG ANATTANANG GANAGAGAGA GANGCCANAG AGATGCATGA	ANGGAANGGA CHANGGGANG ANGRGATTGA NGCTCANGAA AANGCCANAC GGGAANGAGA	OTOGENGAAA AACTITICAGG AAAGTOGAGA TOGTOCOTOTG GACAGCTGGC GAAACTITCCA	AGCCAATAGG AAGGGAAGA AAGAGAAGAA AAATGGGACC TTCCTGAGAC CACCGAAAGT	AANANTOGAG CAACOTGAGT GACCGCCAA GGTCACAGGC ACAGAACCTT TCCCCTGCTA	TCTCCCTTCC TOCTTCGAAG GACTCATTCT TTCCTCCCCAC TTCCACCCCA ACATAGAGTA	GTATTIGCTT TITAGLICCAT ITHGITTICA AFACGAITTA AFALCGAICA GAGIAAITCE	TITICIACATT GAAATGAGGG GCTTGGTTTA AAAAAGACC ITTCCCTCTC CCTGCCCCTA	GAACAACAG TATTAGAAGG TGCCACCATT GGTGCTGCCT TCTCTTCCCA CAGCCTGTAA	CTCAGTGTT TGTACTICAC TGAATTGTGA TGGTTAGAAA CTTCGTGGAT AGTTTGTGGA	AATCATCCAA TTAAACATAC TGCTTAAAAC AGTOTTGCTG TGACTTCAGA GACAAGCCTG	GAAGGGGCAC CITAGGAAGC CCCTTCGCTT CAGTTGCTCG CITCTGGGTG TGCTCCCTTC	GANGGCCCAG ATAAGACAGG GAACACTTGT GAGCACACAG AGCAGCATCT GATGCCCTGT	GOTOTITICAC ATOTICCCCC TOTCTACTCA CCAATCAGTG TOCCATGAGG CCCACGCCAC	CCAAACCTIT CACTITICCAA AGAGCIAGCC GTCCTCCACC CAGTACCATG TCCTAGGCTG	TCTGCATTIG TTAGTGGTAA TATTCTTTAT GTATAATAAA TTTTTATACC CAAAAAAAA	AAAAAAAA ACTCGA	
		S		9	2		15		ç	3		25		20	3		35	

(2) INFORMATION FOR SEQ ID NO: 189:

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 681 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 189:

GGCTACAGGC	TTATCTCGGG	TATAAGCCCT	CIGTATATIT	ATCCCAGGGT	GEARACTIAA
GCTCCCATGT TGCTGGCTGT CCGTACATCA CCCTGTCCCC TGCAGGAGGG GGCTACAGGC	CATCTCCCTC CTGTAGGCCT CTGACTCCCC TCCACTTTTG GOCCCTCAGC TTATCTCGGG	CAGGGGACCA TIGCAGCAIC CICCCCICCT CNGACTCAA GGIGCIGAGG TATAAGCCCT	GGCCCCAGA TCCCTGRTKA CACCTTCCTG GAGAAGACTC TCAAAAGTGA CTGTATATTT	GAGTTCACCA GCAATAACTC CCCACACTCG AAGCAGGTCC AAACCCAAGG ATCCCAGGGT	THEOREMAN
occroroccc	TCCACTITIG	CNGGACTCAA	GAGAAGACTC	ААБСАВЕТСС	CCTTGGGGTC TGTGGGCACTG TCTTCCCAAG ATCCTTCCTG TTGCACAATG GGAAAACAAA
CCGTACATCA	CTGACTCCCC	crecereer	CACCTTCCTG	CCCACACTCG	TCTTCCCAAG
тестевстег	CTGTAGGCCT	TIGCAGCATC	TCCCTGRITKA	GCAATAACTC	TGTGGCACTG
GCTCCCATGT	CATCTCCCTC	CAGGGGACCA	вовссссива	GAGITCACCA	CCTTGGGCTC
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쏬 6 30 25 2 20 5 55 8 5 Ś TATTOGCTAC ACTICIDAAC AACCICAAAG TICAAGAAAG GCAGAACAGA GIITGIACCA ACCACATOTC ACTIOGOGIT TATOGATTIG GITGIGAAGA TICGCIGAAT CACTIGITGA TCTIGITIGA ATATATIGGT GAAAIGGGAA AAGACTACAT TTATGCCGTA ACACCGTTAC GCTGGATGAT TGCTGGGCAA AGGTGGCCTF TTAGAGCTCT TAAAAAGCCCA CAAAAAGGCT GITAGICTAC AAACCATIGC AGCAGCACCA ICAAACCAGA GICIGCCACI TITIGICAIC TOGICICCIC CIGGAIGCIG CICCITGAAT YITITITYIT GAWAAACCYI TIAMAAITAA TATOTTIACA TECCEGAGGG GITTETGEET CETECCEACE CAGGICAGGG TOTGGTCEAG GTCTGTTTCA CCCAGCCCGG AAAGTCAGAG ATGTATATTG GAAAATTTAC AACTCCATCT CCCTAGAGGG CCTGAGAGTT GCTATTOGAC CATGTAGAAT GTTGCAATAT TOTTTACAGG tigaagatec titaategat agagaccite tacacagaca gaceectagt ecagteetac TGAATGAATA CAGAGITCCT GAACTGAATG TICAAAATGG AGIGITAAAA TCGCITICCI CIGTAGCAAT AGCTATIGIT GCAGAAACAT GIICACCCII TACAGTACIC OCIGCCIIAA GCCTCAAGCC ACGCATATGA TAATTYTCTG GAACATTCAA ATTCAGTGTT TCTACAGCCA 2 AAAAAAAAAA AAAAAACTCG A CAGCTIGCTG TGGGGTGCTG ACATGTGTCA CCACTGCCCC CCTTGCCCCC GGGGGGGTCA TOTOYCTOCC TOGOTOGAAT GTOGGCCCCT GCTCCCCOTC AGGTTGTGCT GTCTCTGACC TIGCAGATCA GIGTAGGACI, GGICCATAGG GGAAGAGCTA GGAANTCCAT AGGC RACACCTATA TICGITAIGA ACTIGACIAI AICITAINAT TITAITIGIIW ATTIKGIGKI ACTATOTATO GCCCAATOTR TTTGAGACAT CTCCTCATGT AATTCAGGCA GTTATGGGAG ATTOGTAGAG CCACAGTCAA CACATTTOGT TATATTOCAA AGGCCATTOG CCTCATGATG TAATGCACAS TACTTCACAC CTTAAACTTG CTTTGATTTG GTGATGTAAA CTTTTAAACI ACATTGGTTC CCAGGACGCT CTCATAGCAC ATTACCCAAG AATCTACCAA CGATGATAAG INFORMATION FOR SEQ ID NO: 190: Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 190: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1014 base pairs 1014 960 900 840 780 720 660 600 540 480 420 360 300 240 681 660 600 540 480 60

GAGGAAAAAG ACAGGGGCCT GCTTGCCCAG CCATGCGAGG GATTCCATGC CCACCTGCCC 420

6 35 ဗ ACGOGGICAA TOCCTOCATT CTOCCACTOC TOCAGATCGA CCGGGACTCT GGCAATCCTC AATACACAGA GOOCTCCACA GOTAAGACOT GCCTGATGAA GGCTGTGCTG AACCTTAAGG AGGATOTOGO TOGACTITOCA GAGTACOTGA GOAAGACOAG CAAGTACOTO ACOGACTOGO COGATICIAMA COGATITIGAE CGAGATICOGE TETTICAATIGE OGTETICECOG AATTOGOOO TOAGATAAGA GTOAACOTOA ACTACOGAAA GGGAACAGGT GCCAGTOAGO AGCIGGATITI TOGGAGCGOG CIOCCICCCA TOGAGICACA GIICCAGGOC GAGGACCGG GICCIUGGIG GACCGAGCAG CCICCICCIC CIAGGAIGAC CICACCCICC AGCICICCAG ITITICAGOTT GGAGACATTA GATGGAGGCC AAGAAGATGG CTCTGAGGCG GACAGAGGAA GOTOTCCCCC

> 600 540 480 420 360 300 240 180 120 8

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GCCAACACCG ACGCGCANTG GGAGGAAGAC AGGACCCTTG ACATCTCCAT CTGCACAGAG GAAGCTCCGG GATCCCAGCA GCCGCCACGC CCTGGCCTCA GCCTGCGGGG CTCCAGTCAG 20

CTOCTOTEAG COCCOOCAGE CCCTCCCOGG TTCACTTCCT CCCGCAGCCC CTGCTACTGA

CCAGAACCTO CTTOCTOGAG CTTAGTOCTC AGAGCTGGGG AGGGAGGTTC CGCCGCTCCT

ASTINAGETE CEGYTETECA CEGTGECEGE TEGECEAGGTG GGETGAGGGT GACEGAGAGA

5

TOSCASCAGO STOTOTOCAS ATGSTCASTO TOTOSTOSOT ASCOTOTOCO GACAGOGGAS

ICCCCATTGA GAAGAGGAGW CTGCAGTGTG TGAAGCTCCT GGTGGAGAAT GGGGCCAATC AGCCCCTGGT AMATGCCCAG TGCACAGATG ACTATTACCG AGGCCACAGC GCTCTGCACA 960 900 840 780 720 93

GUAGAACCCA CACCAGCCCG CCAGCCTGCA GGCACTGACT CCCAGGGCAA CACAGTCCTG TGAGCTACCC CTCTYTTTGG CCGCTTGCAC CAAGCAGTGG GATGTGGTAA GCTACCTCCT IGCATIGCCCO GGTCTIOCOGC GCTTCTTCCA GAAGGGCCAA GGGACTTGCT TTTATTYTCGG 1140 1080 1020

COCAACCTOC AGGATCTCAC GCCTCTGAAG CTOOCCGCCA AGGAGGGCAA GATCGAQATT CATIGOCOTAG TGATGATOTO GGACAACTOA GOTGAGAACA TIGOACTIGGT GACCAGCATG TATGATGOGC TOCTOCAAGC TGOGGCCCGC CTCTGCCCTA CCGTGCAGCT TGAGGACATC 1320 1260 1200

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GASTOSTISCT ANGGECCTOT COGGSTGTCG CTGTATGACC TOGCTTCTGT GGACAGCTGT ITICAGGEACA TECTIGEAGEG GGAGITITICA GGACTIGAGEC ACCITITECEG AAAGITICAEC 1440 1380

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WO 98/39448

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INFORMATION FOR SEQ ID NO: 191:

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E SEQUENCE CHARACTERISTICS: (A) LENGTH: 2779 base pairs

SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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9 120 180 240 300 360 420 480 540 80 999

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1500 1560 1620 1680 1800 1860 1920 1980 2040 2100 2160 2220 2280 2340 2400 2460 2520 2580 2640 2700 2760 2779 GAGGAGAACT CAGTGCTGGA GATCATTGCC TITCATTGCA AGAGCCCGGA CCGACACGGA ATGGTCGTTT TGGAGCCCCT GAACAAACTG CTGCAGGGGA AATGGGATGT GCTCATCCCC TACCATCAGO CTACOCTGAA GAAGCAGGOO GOOCCTCACO TGAAAGOGGA GGTTGGAAAC AAGIICIICI TAAACIICCI GIGIAAICIG AICIACAIGI ICAICIICAC COCIGIIGC TOCATIGOTIGO TGACGGGCCA CATOCITIATO CTGOTIAGBGG GGATOTIACOT COTOGTIGGGC CAGCITOTGGT ACTITOTGGG GCGCCACGTG ITCATCTGGA TCTCGTTCAT AGACAGCTAC TITIGAAAICC ICITICCIGIT CCARGCCCTG CICACAGIGG IGICCCARGI GCIGIGITIC CTGGSCATCG AGTGGTACCT GCCCTGCTT GTGTCTGCGC TGGTGCTGGG CTGGCTGAAC CTOCITITACY ATACAGGIGG CITICCAGCAC ACAGGCATCY ACAGTOTCAT GATCCAGAAG CCTGAGCCAG GANNTIGGCG CCCCGAAGCT CCTACAGGCC CCAATGCCAC CAGCCCATGG AGGACAGGA KGAGGAKGGC NACGGGGCCC AGTACAGGG GCCTCCTTGG AGCTCTTCAA AFTCACCATC GGCATGGGCG AGCTGGCCTT CTOCACTICC GOGGLATUGT GCTGCTGCTG CTGCTGGSCT ACGTGCTGCT CACCTACATC CTGCTGCTCA ACATGCTCAT GGCCTCATG AGGGAGGG TCAACAGTGT COCCACTGAC AGCTGGAGCA TCTGGAAGCT GCAGAAAGCC ATCTCTGTCC TGGAGATGGA GAATGGCTAT TGGTGGTGCA GGAAGAAGCA GCGGGCAGGT GTGATGCTGA CCGTTGGCAC GGAAAACTAT GTGCCCGTCC AGCTCCTCCA GTCCAACTGA TGGCCCAGAT GCAGCAGGAG GCCAGAGGAC AGAGCAGAGG ATCTTTCCAA CCACATCTGC TGGCTCTGGG GTCCCAGTGA TAAGCCAGAT GGCAGCCCSG ATGAGCGCTG GTGCTTCAGG GTGGAGGAGS TGAACTGGG AACTETECENG AACKETGTCE TGGETTCCCE TECEAAGGAG GATGAGGATG GTGCCTETGA TTCATGGGAG CAGACGCTGC CTACGCTGTG TGAGGACCCG TCAGGGGCAG AAAAAAAAA AAAAAGGC CCCTGGTGAG AGAGTCAGTG TATCCTOGAA CCAGGARCAG

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INFORMATION FOR SEQ ID NO: 192; ĉ

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LENGTH: 1923 base pairs (A) LENCTH: 1923 base pair (B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLGGY: linear SEQUENCE CHARACTERISTICS: Ξ 55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 192:

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1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 1740 1800 GOGCTGCCGC TOGATCCTGC CCCTGCTCCT ACTCAGCGCC ATCGCCTTCG ACCOGETCOS ETECOCITEOS ETESSECECOS ESCOSOCIOST CAACATGATE CGCTGCGGCC CAGACOTCCT TGGTAATTTA CCCGGTGAAG TACACCCAGA CCTTCACCCCT TCATGCCAAC CSTGCTGTCA AGAGCCTCAT GGAGTACGCG TGGGGTAGAG CAGCGGCTGC CATGCTCTTC TGTGGCTTCA CCTTGGCTGC TGTGTTCCAG ATCATCTCCC GIGCCITICIT CITICISCISC CICCCCAACT ACGAAGATGA CCITICISGGC AATGCCAAGC CACATECTTC CCAGGIACIT CIACACAICI GCCIAACITG GGAATGAATG TGGGAGAAAA TCGCTGCTGC IGAGATGGAC TCCAGAAGAA GAAACTGTTT CTCCAGGCGA CTTTGAACCC ATTITTTGGC agtottcata ttattaaact agtcaaaaat gctaaaataa tttgggagaa aatattttt AAGTAGTGIT ATAGITICAT GIITAICITT TAITATGITT TGIGAAGITG 1GICITITICA CTANTTACCT ATACTATGCC AATATTTCCT TATATCTATC CATAACATTT ATACTACATT ISTAAGAGAA TAIGCACGIG AAACITAACA CITTATAAGG TAAAAAIGAG GITICCAAGA ITTAMIANIC IGAICAAGIT CITGITATIT CCAANIAGAA TGGACTCGGT CIGITAAGGG CTARGGAGAA GAGGAAGATA AGGTTAAAAG TTGTTAATGA CCAAACATTC TAAAAGAAAT AAATGCATAT CATTTOTGAG AATTTCTCAT TAATATCCTG AATCATTCAT TTCAGCTAAG STITATATGI TCAGAACCAG AGTAGACTGG AFTGAAAGAT GGACTGGGTC TAATTTATCA ICHGGCTCT ATCATATAGA CAGGCTTCTG ATAGTTTGCA ACTGTAAGCA GAAACCTACA TATAGITAAA AICCIGGICT TICITGGIAA ACAGAITITIA AAIGICIGAI AIAAAACAIG GCAAAAAAA AGITITATITI CAAGCCITCG AACTATITAA GGAAAGCAAA ATCATITCCI GCTTCATGTT CACTCGATAT GTCATCTAGG AAAGTACTAT TTCATGGTCC AAACCTGTTG TGACTGATAG ATCTGGTTAA GTTGTGTAGT AAAGCATTAG GAGGGTCATT CTTGTCACAA AAGTGCCACT AAAACAGCCT CAGGAGAATA AATGACTTGC TTTTCTAAAT CTCAGGTTTA GGICATTATG ATTITITACC ATTICGACTT ACATAATGAA AACCAATTCA TITTAAATAT CCATAGITGG TAAGGCTITIC CITTAAGIGT GAAATATITA GATGAAATIT TCTCTFTTAA CCACAGGAGA ATTCGGGGAT TTGAGTTTCT CTGAATAGCA TATATATGAT GCATCGGATA AGTICTITAT AGGGTTAGGG TGTGGGAAAA TGCTATATTA ATAAATCTGF AGTGTTTTGI GCTGGCGGC CGCGGCTGGT TGCAGTCTAG CGACCACGGC COCTISTISSTIC GAAATICCTCC CAAGAGGGCG GCGGCAGCGG GTCCTACGAG TOTTOGCCOT CTGTGGACCC CITACATCTA TAACTGGGC TACGGCTTTG GGTGGGCAGC CACGATTATC TCATCCTGGT GATCTGTTTC ATCCTCTCT TCTTCCTGAG AGTGATTGGA GGTCTCCTTG TGGCCTGCGA ACATCATOCC Š 9 13 2 23 8 35 **\$** 45 S 55

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35 8 55 50 45 6 30 25 20 15 5 S CICCCCGGCC CCITITICGCC CCCIGCCIGC CATACIGCGC CIAACICGGT ATTAATCCAA TITTICIAGCA TCCTCTTAAT GIGCAGCAAA AGCAGGCRAC AAAATCICCT GGCTTTACAG ACCGATITIT AAAGIIGGIG CATCTAGAAA GCTITGAAIG CAGAAGCAAA CAAGCTIGAI GENTICATOT CHARGETAGA GECATITIGA ACAACAANIC TACGIAGITA ACITGAAGA GGTAATAGTG ACATGCAGGC ACCTCTTTTA AACAGGCAAA ACAGGAAGGG GGAAAAGGTG AACTGAGGAG AAGCTGATCC AGTTTCCGGA AACAAAATCC TTTTCTCATT TGGGGAGGG ATCCTOGTTC AMACGGGTGC CTGGTCAGAA GGCCAGCCGC CCACTTCCCG TTTCCTCTTT GCCAAAAACA AGACGCGTAC AGCACACACT TCACAAAGCC AAGCCTAGGC CGCCCTGAGC CTCTTCCATT AACCAGTOGC CGGTTGCCAC TCTCCTCCCC TCCCTCAGAG ACACCAAACT AGCTTATTTT GTAAGAGTGA GCTCTGGTGG AGACAAATGA GGTCTATTAC GTGGGTGCCC CCTGCTAGGC GGCCTGCCCA GCTGCCGCCC CCGGACTCTG ATCTCTGTAG TGGCCCCCTC ACATCACCC CAACTCGGGT GCTGCTGGGA ACAGNGCCGG GCCCAAGTCC ATGGAGGTCT GCTCCCACAC CAGCGAGGGC ACCCGCAGCC GCTCGCACAC CAGCGAGGGG GCCCACCTGG AGCOTCATTC TCTOGATOGC ACCCGCAGCC GCTCCCACAC CAGCGAGGGC ACCCGAAGCC CCCCTAGTIGC CCTCGGCTAG CATGACCCGC CTGATGCGWT SCCGCACAGC CTCTGGTTCC CEGICATISCET GECTICATICAL CIRGOLAGICT GECTIGENCIES CITAACCGGET GICTETTIGIT GCTGCAGGG ATCAAGTCTT CTCTGGGGCT GGGCACGTAN AAGAGCATGT GGCTGGTGGA GECCTIVECA TCTCTCTCTC CGAGTGGACA TOGAGAGGAC GGGGGCCCAG CAGCTGGATG AGGCTCAGGG GGACACTCTC AAAATTACAC AGCTTTTAAC AGGTGGCAGA ATTGGGGTTC Ą AGACCCAGAT CTGGGTTCAA GTCACTCATG GTGTGATTGC GGCATTCCTT CCCGCATCTG (2) INFORMATION FOR SEQ ID NO: 193: CCAATAAACC AGGTATTCTA AAAAAAAAAA AAAAAAACTN GAGGGGGGGC CCGGTACCCA CACATTATTA TITTICTAAGT TOTOGAAAAA GCTAATIGTA GITTICATTA TGAAGITTIC Ĕ E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 193: ô (B (D) TOPOLOGY: linear (A) LENGTH: 2346 base pairs TYPE: nucleic acid STRANDEDNESS: double 1200 1260 1080 1020 1923 1920 900 840 720 660 60 540 1860 780 480 420 360 300 240 180 120

> 8 35 30 25 20 2 5 S ACAATC GOCCTTCCCT GOCCTTGAGT AGAAAAGTCG GOGATCGGGG CAAGAGAGGCC TGAGTACGGA CCTICCCICT CCTGCTTCCT CTTTTCCTGC TCCCTAACCT TTCCCCGAAT GGGGCAGCAC GOCGGAATTG GGGTTACTCG ATGTAAGGGA TICCTTGTTG TIGTGTTGAG ATCCAGTGCA AAAAAAAAA AAAAAAAACT CEAGGGGGGC CCGTACCCAA TTCGCCGTAT ATGATCGTAA TITICCAGACC AATAAATTIG TAACTITIGCA AAAAAAAAA AAAAAAAAAA AAAAAAAAAAA TICLAGICITIC CAGGIGGIGT GAGAGGICAAT GACTICGTTAC CTGCCGCCCA TICACCTTGGA GTANGICAGO CACTOGGACO CGAGGATTTO TOGGACCCCG CAGTTGGGAG GAGGAAGTAG TICATITIGG CICACCGIGG ATITICICAT AGGAAGTITG GICAGAGIGA ATIGAATATI TOGGAAACTA TIOTOCACAA GICTITICCAG AGGAGITTICI TAATGAGATA TIITGTATITIA CACTGACGIT TCTGGGCGGC CAGTGCGGCT GCCAGGTTCC TGTACTACTG CCTTGTACTT COTGOGGAGC GOTGOCTGAG AAAANOTAAG GATTCTOGAA TACATATTCC ANGOGACTTT COGTICACOTT CITICGATICCT CAGAACTICTT TOCTICTICTC OGGGTOGGGG TGGGAACTICA ADTITICAAT CITCGACAGC TOGOCTOGAA COTGAACTCA GTAGCTGAAC CIGICIGACC CAGCTOCCCT OCCIVITATIC ACTOTCCTTG CAGGGGCCGG ACTAGGAGCA CTGGGGTGGG TICAGGGAGA AGAICATTIA GATTIGITIT GCATICCITA GAAIGGAGGG CAACATICCA TOCCTOTOCT COACAGECCC AACCTCCCAC CCCTGATACA TGAGCCAGTG ATTATTCTTG OTTOTIGATIT CTOTOGRATICE CAGCITOGIT CCAGGRAFIT TOTOTIGATIG GCTTAAAICC ACAAAAATAT TTCAGCAAAC GTIGGOCATC ATOGTTTTTIG AAGGCITTAG TICTGCTTTC 2340 2280 2220 2160 2100 2040 1980 1920 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320

(2) INFORMATION FOR SEQ ID NO: 194:

Ξ SEQUENCE CHARACTERISTICS: (A) LENOTH: 3054 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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55 ACCICMAGOT OCCCITOTOGN ACGAGCAACT GGACTATAGC AGGGCTGGGC ICTGTCTTCC TATICTICANCE ACCEPTATE CTACATATION TAGGENGENE TIGANATATICE TANCECCETA AAAGGAATAG GTAGGAGACC TCTTCTATCT AATCCTTAAA AGCATAATGT TGAACATTCA TOGTCATAGO CICACICITI COCCCAAATO TICCICIOGA CCITIGGAGO CAAGGIGGTA 240 180 120 6

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 194;

	WO 98/39448 P	PCT/US98/04493	W	WO 98/39448	PCT/US98/04493
	425			426	
	TICAACAGCT GATGCCCTAT AACCCTGCC TGGATTTCTT CCTATTAGGA TATAAGAAGT	300		GACAAAGAAG GAAAAGAGTA TCAAAGGCAG AAAGGAGATC ATTTAGTTGG GTCTGAAAGG	2100
	AGCAAGATCT TTACATAATT CAGAGTGGTT TCATTGCCTT CCTACCCTCT CTAATGGCCC	360		AAAAGICITI GCIATCCGAC ATGIACTGCT AGIACCTGIA AGCATTTIAG GICCCAGAAT	2160
ς.	CTCCATTTAL TTGACTAAAG CATCACACAG TGGCACTAGC ATTATACCAA GAGTATGAGA	420	2	GGAAAAAAA ATCAGCTATT GGTAATATAA TAATGTCCTT TCCCTGGAGT CAGTTTTTT	2220
	AATACAOTGC TITATGGCTC TAACATTACT GCCTTCAOTA TCAAGGCTGC CTGGAGAAAG	480		AAAAAGTIAA CICITAGIII TIACITGIII AATICIAAAA GAGAAGGGAG CIGAGGCCAI	2280
2	GATGGCAGCC TCAGGGCTTC CTTATGTCCT CCACCACAAG AGCTCCTTGA TGAAGGTCAT	. 095	9	TCCCTGTAGG AGTAAAGATA AAAGGATAGG AAAGATTCA AAGCTCTAAT AGAGTCACAG	2340
2	CITITICCC TAICCIGING INCCCTCCC CGCICCIAAT GGNACGIGGG TACCCAGGCT	009	2	CTITCCCAGG TATAAAACCI AAAATTAAGA AGTACAATAA GCAGAGGTGG AAAATGATCT	2400
	GETTETTGGG CTAGGTAGTG GGGACCAAGT TCAITACCTC CCTATCAGTT CTAGCATAGT	099		AGTICCTGAT AGCTACCCAC AGAGCAAGTG ATTTATAAAT TIGAAATCCA AACTACTTTC	2460
15	AAACTAGGST ACCAGTGSTA GTGGGAAGAG CTGGGTTTTC CTAGTATACC CACTGCATCC	720	15	TIBATATCAC TITIGATCICC ATTITICCCA GACAGGAAA TAIGICCCC CCIAACTITIC	2520
	TACTCCTACC TOSTCAACCC OCTOCTTCCA OSTATGOGAC CTOCTAAGTS TOGAATTACC	780		TTGCTTCAAA AATTAAAATC CAGCATCCCA AGATCATTCT ACAAGTAATT TTGCACAGAC	2580
ξ	TGATAAGGA GAGGGAAATA CAAGGAGGC CTCTGGTGTT CCTGGCCTCA GCCAGTGCC	840	00	ATCTCCTCAC CCCAGTGCCT GTCTGGAGCT CACCCAAGGT CACCAAACAA CTTGGTTGTG	2640
3	CACAAGCCAT AAACCAATAA AACAAGAATA CTGAGTCAGT TITTITATCTG GGTTCTCTTC	006	3	AACCINACTO CCTTIAACCTT CTGGGGAGG GGGATTIAGCT AGACTIAGGAG ACCAGNAGTG	2700
	ATTCCCACTG CACTTGGTGC TGCTTTGGCT GACTGGGAAC ACCCCATAAC TACAGAGTCT	096		aatgogaaag gotgaggact teacaatott goectoteag agettgatta gaageeaaga	2760
25	GACAGGAAGA CTGGAGACTG ATCCACTTCTA GCTGGGAACT TACTGTGTAA ATAAACTTTC	1020	25	CAGTGGCAGC AAAGGAAGAC FTGGCCCAGG AAAAACCTGT GGGTTGTGCT AATTTCTGTC	2820
	AGAACTIGCTA CCATGAAGTG AAAATGCCAC ATTITIGCTTT ATAAITTTCTA CCCATGTTGG	1080		CHGAAANTAG GOTGGACAGA AGCTTOTGGG GTGCATGGAG GAATTOGGAC CTGGTTATOT	2880
30	GAAAAACTIGG CTTTTTCCCA GCCTTTCCA GGGCATAAAA CTCAACCCCT TCGATAGCAA	1140	9	TOTTATTCTC GGACTGGAA TTTTGGTGAT GTAAAACAGA ATATTCTGTA AACCTAATGT	2940
3	GTCCCATCAG CCTATTATTT TITTAAAGAA AACTTGCACT TGTTTTTCTT TITACAGTTA	1200	3	ctgtataaat aatgagggtt aacacagtaa aatattcaat aagaagtcaa aaaaaaaaa	3000
	CTTCCTTCCT GCCCCAAAAT TATAAACTCT AAGTGTAAAA AAAAGTCTTA ACAACAGCTT	1260		aaaaactog aggogggcc cogtacocaa titinccaaat agagathota teac	3054
35	CTIGCTIGIA AAATAIGIA TIAIACAICI GIAITITIAA AITCIGCICC IGAAAAAIGA	1320	35		
	CTGTCCCATT CTCCACTCAC TCCATTTGGG GCCTTTCCCA TTGGTCTGCA TGTCTTTTAT	1380		(2) INFORMATION FOR SEO ID NO: 195:	
40	CATTGCAGGC CAGTGGACAG AGGGAGAAGG GAGAACAGGG GTCGCCAACA CTTGTGTTGC	1440	64	(i) SECTION CHARACTERISTICS	
?	TITICITGACTIG ATCCTGAACA AGAAAGAGTA ACACTGAGGG GCTCGCTCCC ATGCACAACT	1500	?	(A) LENGTH: 907 base pairs (A) TUDE: mileic anid	
	CTCCAAAACA CTTATCCTCC TGCAAGAGTG GGCTTTCCAG GGTCTTTACT GGGAAGCAGT	1560			
45	TAAGCCCCCT CCTCACCCCT TCCTTTTTC TTTCTTTACT CCTTTGGCTT CAAAGGATTT	1620	45	(xi) SEQUENCE DESCRIPTION: SEO ID NO: 195:	
	TOGADAGAA ACADTATOCT TTACACTCAT TTTCAATTTC TAAATTTGCA GGGGATACTC	1680	Ū	SCHARGER GROSCORUM CITITATICA CONCENSION CANCERS CONCENSIONS	09
20	AAAAHAGGG CAGGIGGCCT AAGGCIGCIG TAAAGIIGAG GGGAGAGGAA AICTIAAGAI	1740	55	CCCTGGCCTG GGCCTGCACC AGCCTGCGNG CGGGCTCCCA CAGCAGCCCC CTTCCAAGCA	120
	TACANGATAA AAAAGGAATC CCCTAAACAA AAAGAACAAT AGAACTGGTC TTCCATTTTG	1800	J		180
	CCACCTTICC TOTICATIGAC AGCTACTAAC CTGGAGACAG TAACATTICA TTAACCAAAG	1860	·	COSANGOCITY TOTOCHOOC PACABOCITES STUDANCIACES TOCCAGOCITY TOTOCHOOC	240
55	AAAGTOGGTC ACCTGACCTC TGAAGAGCTG AGTACTCAGG CCACTCCAAT CACCCTAGAA	1920	55	отнастью населена самента предоставляние	
	GATGCCAAGG AGGTCCCAGG AAGTCCAGGT CCTTAAACTG ACGCTAGNCA ATAAACCTGG	1980		CONTRACTOR CANADATA C	
09	GCAAGTGAGG CAAGAGAAAT GAGGAAGAAT CCATCTGTGA GGTGACAGGC AAGGATGAAA	2040	9	GOACHGANA CHGCHGCTC TGTAAGATG GTGCTGCGCT TCCGCCTCGG GGGCCTAGCC	420
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25 20 30 15 5 S CTYTOGSCOG AGGGCCTOGC TGATGGGTTG GGGCCACATC CCTGCTTCTT GCTCCTGCTC AACAATO GGCACGACGA GCGACAGGGA GTGGGCAAGG GGAAGAAGCA GCTTATTTGA CTAACCAGCC aaaaaaaaa aaaaaaactg gagggggggc ccggtaccca aatcgccgga tatgatcgte GAGTOGCTGG AGTOGTCAAT AAAGCCACAT GTGCCTOTOG CCCAAAAAAA AAAAAAAAAA ANCONONIO CONTROCCON NOVONACONO GACCIAGOAC COAGCACONT NOVONACON CACCACAGIC ACCITICACIO GGAATGATOC GCTGCAGCCA GCTGGCCCCC AGGGCCTTGC AATCTTGAGA GGGTCAGCCT TOCTGAGCCT ATGTCTGCAG CACTTCTTGG GARGCCTGGT (2) INFORMATION FOR SEQ ID NO: 196: ACCIONACACA CTACACICTY CYBOCIONOC YBGAGITOCY GOGGAAGITG CYBITOGOCA TOTCAGACTO CCTTGGTCTT CCACCTTGGA CACCCTGGGG GCCAGCATGG ACGCTGGGAC Ê Ě SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 196: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double 9 TOPOLOGY: linear LENGTH: 1290 base pairs 900 840 907 780 720 660 600 540 480

CCTCTOTOGT CCACCAGCOT CTTGGCTTGG TGGGAGGGCT CTCAATCAGC AGGGCCCCAG 120

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TOTOGRACICA COCCTOCTOC AGGCOATGCT COTTGRACTT GGARATGTOR ACCOGRACICO KAGGGCAAGA AGAAGTOGGG CAAAGCCTOG CGCTCOGCCG CGGTCGCGGC AGCTTTOCAA 240 180

TIMACACCAG CCCICCAGCA ICIMATAGAC TIGAAICIAC ICIMAACGAA TAITIMAICC AACCTCAACT ACATTGTAGC TCAGTCCAAC GACTAACCCT GAAATGGGGG TGTTCCAGCC 360 8

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45 TICADCGAGA TGGCCAAGCG GICCCCIGGG GCCIGIGGCA GCGGGCTIAI CCITCICIGI TOCCAACCIT GOOGTOCGAC CTCCTCCGOC CCCATGOGGT GACCCCGTCC GTGTCTGTGT 480 420

S CTGTCCATAC GTGTGAGTCC AGCTAAAAAAG ACAAAAACAGA ACCCGTGGGC CCAGCTCGG AGGTGCGTGG AGAAGGCTCC GACGTCTCCG AAGTGCAGCC CTTGGGATGG CATTCCGTTC

55 TGTGCCTTAT TCÇTGGAGAA TCTGTATACG GCTCGCCTAT AAGAAATATA GCCTCTTCAI

GOTACCCAAT TCGCCCAATA GTGAGTCGTA TTACAATTCA CTGGGCCGTC STTTTAACAA

35 ၓ GOTOTOCCTG GATOGTCOTG TAGGTGAGTT TTACCAAGGA TTATGGTAAC AAATGAGTGA SCITTIGGAGO ATTIGIATIG ACCITITIACA GIATICATIT TICAACICAA GGCAATGGCI GACCTUTATO GAGAAAATAT TGAAGNINCAT TAAAGAAGAC CTCATANTAG GAGAGAATOT TTCTACACCA ACTCTAATCC ATAAACGGGT CTTATGACAT CTATGAAGTA GTAGCAAGAC 240 180 120 8

ATTATOTAGG TAGAAAAAA TOCAAGCAAG CTOTTAAAGA TOTTOGATOO CATTATATAG THIGINIAGC TGARATCIGI ANTICARICA CITTITICICI TITRICCICI ARCCARARA AUGCITAGIG IGIATITICIC ICITIGAGAC ACIGIAATIT CIACCAGAAA ITICCAGAGC 420 360 8

TIGITIAATT TIGCATCCCA AATGITTITIA ATCITIGIAT ATTITTIAAA AAYCCTITIC 480

TOCTOATOAT TECCTTTTTT GIEGITIGTAA ATAGACTTAC TIGGACTITE AAGATGAGIT 540

5

8

ACTOCTTOTO ATOTTACAAA TATOTOATAT GOTAATTTTO ATAACAGATG TOAOTYTTGA 600 9

600 540

COTEGTIGAAC TOGGAAAAACC CTGGCGTTTA CCCAACTTAA TCGCCTTGCA GCACATCCCC

840 780 720 660

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Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 197: 25

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INFORMATION FOR SEQ ID NO:

197

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1020 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

12

AGGGGANCCC CCCATTIAAA ATTITNGGIN

CCAANCAATT TITITIOGOGG GICCCNAAAG GICCCCCTAA AANCITITIT CGGAACCCNA AAAGGGGAAA AAACSYTTTT YTGGGGGGGAA ANGGGGGCCCC CNTACTTTNA ACAYCCCCCC TICCCCAYIT TIGGAAACAA AMTYCCCCCT TITTAAAAAA GIIGGAACCC CCAMCCYTCC

1260 1290

1200 1140 1080 1020

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780 720

ACCAMBANT GGIGATITOT TTATAMGAAA AAAACTGGCT TCATTTCTGT GAAATTGCTC TICAAIGAIG CITACCAICI ATITIAGCCA CIGAGCCIII TAITAITIOI CIATITOTAA PTIGAAAATT ICTTITTACA OSIGTAAGCC AACTGAGATA COGTGATGGT GTIGATTICI

AGTITÄTTIG ICTTAACICA TITAATAAAT ATACIGITIA ICIGITICIG AAIGGOGACI

840

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GAACTITITIG GATATIGATA TIGATITIGAA AATATITITIGG AATTITITITCT ACTIGAAATI

TTAGAAATCT AATKGAAAAT TCTATAATGT ACTGAAAGTA WGGTTGTGTA CAGTGAKCAC 960 900

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TOCKOGITWA AWIITITIGIT TAAAICARCT CAATITITIT AACOCAATAA GSCCGAAATO

TTTGCGCAGC CCTGAATGGC GAAATGGCAA ATTGTAAGCG TTTAATATTT TKKTTAAAAT

960

900

CTITICOCCAG CIGOCGITIAA TAGCGAAAAA NGCCCGCACC CGAATCGCCC TICCCAACAG

COCCADATICE CEYTTATTAA TICCAAAAAA ATAAACESAA AAWGGGTTIG AATTITITIKT

_	WO 98/39448	PCT/US98/04493	W0 98		PCT/US98/04493
	429			430	
	TCTCTAATAA TATGATGACT TGCCCTAAAN GAGGAGGGAC ATGTCCCACT TTCCACCACG	1020	5	(2) INFORMATION FOR SEQ ID NO: 200:	
2			5	U	
	(2) INFORMATION FOR SEQ ID NO: 198:				
٠	(1) SEQUENCE CHARACTERISTICS:		•	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	•
2	€ €		<u>e</u>	(xi) SEGIENCE DESCRIPTION: SEC ID NO. 200.	
	(C) STRANDEDRESS: double		į	יחם יחול משתור מחור ביותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יום יותר יותר יותר יותר יותר יותר יותר יותר	
	(D) TOPOLOGY: linear		8	CCAGGGAAGC CCCARGCCTG TCCTGAATTG ACATCAGTGC TTCCCTGAAC TGCCTCCCC	09
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 198:		15 AC	ACCCTIGGC ATTAICCCAG GAAACTTAIG TTTTCTAGAA GCIAAGCAGC TGCTGGGACT	120
	ANTICOCGAA GCTGAGGGTT GTGTGCCATC GGGGAGCCA AGTCTTTTGA CCGGAGCCTT	09	ฮี	CAGGGACTIGG TIGCAGGTAGG CTGAGTGGCA GCTCAGTCCT AGAAGGTCTC TGAAGATCTG	180
ę	CCCGGCGCAG AAGANCTGAA GITGALTITGA GAGCCTGTKT TIGGGGTTRA GCCGAGCTGC	120		GACTGAGGAC CYTOCTACTC CCCAAGCCAG AGCCCATCAG CCAGGCCTGC TGTGAGCCAC	240
3	TECHNOCITY OTCHCONIC AGAINAGA YARTTICLAA CONGOCIOG COTGECTTAL	180	_	CTOCCTOTICG ACTOCTGAGG TCAACCAAAG GCTGGCAAGC TCTGGGCCTC ATTTAAGGGA	300
	GATOTITICATE AACCCAGGGG GGGCTTCTGC CCTCTACTCG TGCCAGGCCC ACTTGCCAGG	240	Ē	ITCTGATGAG CCGATGGGGC CTGGAGGCAG CCCATTAAAG CATCTGGCTC GITTITTGGAA	360
25	CAGARCOCT COCCAACCT TCAGGCTGC TCGGAGTCAC CTGTTGGAAT GGACTAAAAG	300	25 AM	алалалала алалас	376
	GACCETTOTG TOGGAACAGG TOCTCCAAAC ACCCTOCTGC TOGCTGCCAG GCAGGCCCTC	360			-
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Š.	CCCGAGCCCA AGCATTIGGC TCTGCTTGCC CCAAGGGGAC AGGAAGCCTC TTGGGCCTCT	084		(4) INFORMATION FOR SEQ ID NO: 201:	
	receptedte gacagoge ectocoptus ecteadataa acte	\$25 \$25 \$25		(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1192 base pairs	
35		E	35	(B) TYPE: nucleic acid (C) STRANDENESS: double (D) TOPOLOGY: linear	
	(2) INFORMATION POR SEQ ID NO: 199:			(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 201:	-
40	(i) SEQUENCE CHARACTERISTICS:	4	λο σα	CCCAGIATAT TECTATAACA TITIATETIAG TGAACTIATA AIGITECTIT GIATTAAATT	09
			AT	AITAGAITAT ATCTITAGAT AATAITGITA CINAATTAGF AGGIAATATA TAITITPAITC	120
ţ	(C) STRANDEZNESS: double (D) TOPOLOGY: linear	4	45 45	AMAMINAMI TOTICCATCTA ATOTICTACCA ATTANTOTAC TIGTAGATGT ATCITATCTT	180
5	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 199:	r		AUCTICAGIC TITIGCIGCCC CIVATCAGOT GICAAGAACT CITCUCCCCT GOGGAAGTIT	240
	GTGATACAAG GAAGGGTGAT CATCATGTT CACCATGCAA TTCCTGGTCA CAGCCTTTCT	. 09	£	ITCTITITICA GGAGGAGGA GGCTITICCC AGGIAAIGIG ICTAGAGTIGI TGGGCAGAAR	300
20	OTHORNOCIA CITICIDACIO TITIGIANIGI COCCADATOC CIAGRICITOT COCCEDICIA	120 5	50 AAT	ANICTOGGAC CACACCACAC CAGITICICC CITAATCCAC GICATITIGCC TICIAICCCA	360
	GAAGGGCTTC TTCATAGATT AGAAAATAAG AATGAGTGAC ATTTCCTATG TGCATATAAG	. 180	g	GCTATGTTTC CAGTGTCCTC TGGGTGTTTC CAACAGCAAC AACAAAYGAA TAAATCTCTG	420
ď	AAGGAGCCAC AAGACATGTC TITTAAATAA AAGGACAGTG TCCATCCTTT TAACTGCCGA	240	55 KTG	KIGAGITGIT IMITIGITCI ICACITIGIT TIACACIGIA MITICIGAGI TIMIGGGIGI	480
r r	ATAGAACCTT GOTCTCATCC TCCTGGAGCT AGGSCTTAAA ACAGCTTCTG TGTTTCTSAT	300		ctgtgaatta aaaaggaaaa gtrgaabtaa gtaaaactca gcttgaagga aaitatacata	. 540
	TKGICTCART GITTIGCCAA GGITTIAITC GG	332	AAT	AATAAGATAA AGCTGACCTG TAGATATARR CAGGTTATAA RAGCTTAGAG TTGTCTAAGT	. 009
9			60 TGR	TGRGTGCAAA KTTTCCTCTG ATCTTTCTGA TGCCGARACA AAAAAGGCAG TCATGTTTGT	. 023

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S COTABIOTAT GCTCCTTACT CABAGAGIGT GGTCCCABAC AGCCTTTOGG AGGTCCTCCT CCITATITCT TCTAAACICA CCATTAATCT GAATAATAGT CAAATITAGG GG AGTCATATTT TCTGGGAAAT ATTTCCAGTG TTTATTTGCA CTTTAGCCCA CTCTGTGTAG ACTOTOGACA COTTITIOTI AGITTATOTO TACATGCAGG GTGTGCAGCA GCOTGTTCAA TGATTCATGG ATGAAACCTG GAACATCTTG AGGACTGAGT TAACCATAGG TCCTTAAATA CTANAGENEG GEGTGTTGTG CAGEGGAMAT GGTCATETGE TGCTANAACA CAGETTECAT TICACACCGG CCIGGCAGTA AACACTIGTA GIGTIGIGCA GIGGAAAACGG ICAICTICCG TOYGTGCACC AAGTCTKAAC CACCACCTTC ATGGGACATA GRTTATOTOC TGGAACATAT WATGTGATTG GAATGGAACC CGARAAGAGA GCAYGCTGTG TTCTTGGGGA CAGGAAAGCT 1192 1140 1080 1020 960 900 840 780

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(2) INFORMATION FOR SEQ ID NO: 202:

ATGACCTOCA GCAGAATCTO AGCAGCTCAC ACCGGGCCCT GGAGAAACAG ATTGACAGGC TOCCOGGGAA GCTOGATOCC CTGACTGAGC TGCTTAGCAC TGCCCTGGGG CCGAGCAGCT

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 203 (C) STRANDEDNESS: double (D) TOPOLOGY: linear

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WO 98/39448

(A) LENGTH: 847 base pairs (B) TYPE: nucleic acid

(1) SEQUENCE CHARACTERISTICS:

8

(2) INFORMATION FOR SEQ ID NO: 203;

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50 AGGTGGAAGA AATCCCTGAG ACACCTTGTG AAAGTCAAGG AGAGGAACTC AAAGAAGAA

ટ COGGGAGICA GGCAGTICCG TCACCAGCTA CICCATICTOA GGCACTITICT AGTGTGTTAG

GITIGCAAGOG CAGAGAAGAA ACGGTAGCAG AAGATGITTIG TAITIGATCTC ACTTOTGAIT

ATGACAAAAC AAAGGGAGAT GATACAGACA COMGGGATGA CATTAGTATT TTAGCCACTG

AAAATGATAG TATCCTGATG AATCCAGCAC AGGATGGTGA AGTACAACTG AGTCAGAATG

CACAGATTGA GGATACGGAA CCCATGTCTC CAGTTCTCAA TICTAAATTT GTTCCTGCTG

GTCAGTCCCC AAAGATOGAG AGCTTGAGTT CTCACAGAAT TGATGAAGAT GGAGAAAACA

ATCTTGGGCT ATCTTTGACA GGGGATTCTT GCAAGTTGAT GCTTTCTACA AGTGAATATA

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ATCAGGAGGA AGCTATOGAA ATTAAAGAAC ACCATCCAGA GGAGGGGTCT TCAGGGTCTO

ATATOGAGAG TOTTCCOTTG CACCITTCTC TGACTGAAAC TCAGTCCCA

589 540 480 420 360 300 240

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:

50

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(2) INFORMATION FOR SEQ ID NO: 204:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 852 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 202:

(A) LENGTH: 589 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

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AGGACACCAT TITITCCAGAG CIGCAGAGAG CACCIGOTOG GGAGGAAGAA GIOTAACICA

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AGAACTGGGT ATGAGGCTGG GGCGGGGCTG GAGGTGGCGC CCCCTGGTGG GACAACAAAG

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GOSTACTAGO GOCCCOGATC CAGGATTCTG GGAGGCTTCA GTTACCGCTG GCCGAGCTGA

CCAGCTCTCA GTTACAAGTG CAGGCGACTG GAGGCAGGAC TCTTGGGTCC CTGGGAAAGA

TCACCCCCAC TCTGCATACC CTCATCAAAA ACACTCTCAC TATGCTGCTA TGGACGACCT

20

GOOGLACOCTO GCTANAGTOG GGAGGCCTTG GCCCACCTGA GGCCCCAGGT GOGAACATGG

CAMPIOCAMO GACCAMAGOS GOCCIGOCIT OGATIOGOTIG GCTIOCIGAT OGCIGCIGGA

CCCAGTACTS AGTOGTISTAC ATCOTCTCTG CCACTCCTGA CCAGCCTGAA CAAAGCACCT

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TOCAGAACCO AGCOAGCAGT COAAGTAGCT GGACCCACGA GGAGGAACCA GGCTACTTTC

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GGAAGCTCCG GGAACAAGTG AACTCCATGG TGGACATCTC CAAGATGCAC ATGATCCTGT JOCACGAGCG CAAGCTOCTG GCCGCCATCA ACGCGTTCCG CCAGGTGCGG CTGAAACACC

432

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AGCTCACACC GGGCCCTGGA GAAACAGATT GACACGCTGG

COOGGAAGCT GGATGCCCTG

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TOCATEGTOG ACATOTOCAA GATGCACATG ATOCTGTATG ACCTGCAGCA GAATOTGAGO GCCATCAACG CGTTCCGCCA GGTGCGGCTG AAACACCGGA AGCTCCGGGA ACAAGTGAAC ACAAMCATAC TOOCHOGAAG GAGTCTCATG CTGCCCGCAG CATCAGCGCA ACNCNTGGCC

ANOTAGETOG ACCCACGNAG GAGGAACCAG GCTACTTTCC CCAGTACTGA GGTGGTGGAC ACTIGAGETICS TETAGEACTICS SCHOOGGECG AGGEAGETTS CAGAACCCAG SCAGEAGTES

360

AGCCTTGGC COACCTGAGG CCCAAGGTGG GAACAITGGTC ACCCCAACTC TCCATACCCCT	CANCADADAD ACTICICATES SECTIONARIO GACCACCIÓN ACTICIDADES PACADOSAS	S CONTRACTOR CONTRACTO	GONTICHIGGS AGCITICAGT PACCOCTICS CHARTERIANS AACHTGOSTAN GACTATAGAS	10 cococcrisca corcoccocc corcorroga chachangas cachachtri mychanacht		Designation of the second seco	I INTERNATION INCOME AND AND AND AND AND AND AND AND AND AND	AMILLOMIC TOCCTAINET AMENOCCERN ANAN	20 (2) INPORMATION POR SEQ ID NO: 206:	CONTRACTOR CONTRACTOR	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1130 Base pairs (A) ENTYPE: nucleic acid (C) STRANDENESS: double (D) TOPOLOGY: linear	30 mytysagar chargonas coveryme metalosas amenance commences	TOUCHOUSE CHARACTERS SCIENTIFIED STORMS THE STORMS THE STORMS STORMS THE STORMS	35 GACHGGARAA ACAAAANGKO CAAFAACKWO AATTITTAKCT TAGAGATCKO TICCAGCCTAR		TABLETHORIZE THE STATE OF THE S	40 CANTISTICATO ANTICATOR ANTICATOR TOCOGCIETY TECHNISTICATOR ANTICIDATES	GCAGAAAGGA TGGCCCTGAT GCAGCAGCAG CGCCAGCTCT ANATAAAAA TAATTCACAC	45 титеменсти селаноселет асадетовал завлеенея даластален итееланеет	TICATCITAC AGGIGAACAA ACTOTGATGA TGCACATGTA TGTGTTTTGT AAGCTGTGAG		50 COAGGCACH TGACTCCCAG TCTGGTCCC TGTCTACACC AGACAACAA GCACACCAG	CACACHTICCC TCACCTCTT ABCABACTT CTTCABCAGA BACTCTTTAC BARCTCTT	55 TECHNOLOGIAN TRANSPORCE ACTIVITIES ACTIVITIES ATTENDED ATTENDED		יייייייייייייייייייייייייייייייייייייי	CLUMBANGCA AIGITITIAC ITTATICCATA AITCATTOTT GCCAAGGAAT AAAGTGAAGA
920	089	240	009		720	780	940	852					09	120	180	240	300	360	420	480	540	009	099	720	780	840	006
ATNOTICITY TOCCACTICGN TONGCCAGE CCTGAACAAA GCACCTCUAG TGCAAGGACC	AAAGGGGCC CTGGCTTGGA GTGGGTTGGC TTGCTGATGG CTGCTGGAGG GGACGCTGGC	TANAGTIGGGK AGGCCTTGGC CCACCTGAGG CCCCAGGTGG GAACATGGTC ACCCCCACTC	TIGNIACCT CATCANANC ACTOTONIA TACTIGNIATO GACACOTOC AGOTOTICAL	TACAAGTOCA GGCGACTGGA GGCAGGACTC CTGGGTOCCT GGGAAAGAGG GTACTAGGGG	CCCGGATCCA GGATTCTGGG AGGCTTCAGT TACCGCTGGC CGAGCTGAAG AACTGGGTAT	сменствное свенесть ватенское сстентная смемлины влемскитт	TTCCAGAGCT GCAGAGACA CCTGGTGGGG AGGAAGAAGT GTAACTCACC AGCCTCTGCT	CTTATCTTTG TA		(2) INFORMATION FOR SEQ ID NO: 205:	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1354 base pairs (B) TYPE: nucleic acid (C) STRANDERNESS: double (D) TOPOLOGY: linear	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 205:	GATTCGCCAC GAGGCTTGCT GGAGCAGGAG AAGTCTCTRG CCGGCTGGGC ACTGGTGCTG	GCASGARCTO GCATTGGACT CATGGTGCTG CATGCAGAGA TGCTGTGGTT CGGGGGGTGC	TOGGETGTCA ATGCCACTOG GCACCTTTCA GACACACTTT GOCTGATCCC CATCACATTC	CTGACCATOG GCTATGGTGA CGTGGTGCCG GGCACCATGT GGGGAAAAT CGTYTGCCTG	TGCACTGGAG TCATGGGTGT CTGCTGCACA GCCCTGCTGG TGGCCGTGGT GGCCCGGAAG	CTOGRAPITIA ACANGGCAGA GAAGCACOTG CACAACTTCA TGATGGATAT CCAGTATACC	AAAGAGATGA AGGAGTCCGC TGCCCGAGTG CTACAAGAAG CCTGGATGTT CTACAAACAT	ACTCOCAGA AGGAGTETCA TOCTGCCCGC AGGCATCAGC GCAANCTGCT GGCCGCCATC	AACGOCITICE GCCHGSTGCG GCTGAAACAC CGGAAGCTCC GGGAACAAGT GAACTCCATG	STOGACATICT CCAAGATIGCA CATGATCCTG TATGACCTIGC AGCAGAATCT GAGCAGCTCA	CACCOGGCC TGGAGAAACA GATTGACACG CTGGCGGGGA AGCTGGATGC CCTGACTGAG	CTGCTTAGCA CTGCCTGGG GCCGAGGCAG CTTCCAGAAC CCAGCCAGCA GTCCAAGTAG	CTGGACCCAC GAGGAGGAAC CAGGCTACTT TCCCCAGTAC TGAGGTGGTG GACATCGTCT	CTGCCACTCC TGANCCCAGC CCTGAACAAA GCACCTCAAG TGCAAGGACC AAAGGGGGCC	CTGGCTTGGA GTGGGTTGGC TTGCTGATGG CTGCTGGAGG GGACGCTGGC TAAAGTGGGK
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AACAGCACCT TTTAATATAT AGGTCTCTCT GGAAGAGACC TAAATTAGAA AGAGAAAACT 1020 960

GTGACAATTI TCATATTCTC ATICTTAAAA AACACTAATC TTAACTAACA AAAGTTCTTT

TCAGAATAAG TTACACACAA TGGCCACAGC AGTTTGTCTT TAATAGTATA GTGCCTATAC

CATGAGACAG GATTATAGTO CCTTAACCGA TATATTTTGT GACTTAAAAA ATACATTTAA

AACTOCICIT CIGCICIAGI ACCAIGCITA GIGCAAAIGA TIAITICIAI GIACAACIGA

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1140 1080

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 208: (A) LENGTH: 697 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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E SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 208:

CAGTETTEAG CTCNAGGGTT TTAAAA GOCCTITIGCC CCITAGRAAA GTAGCTITTA GGGGCAAAGA TITIGTIGATI TICCCCATTA

> 1166 1140 1080 1020

TIGGGCITTI ANGGITCAGA GACIGIGGGC TIGGGCACCT GCGCCCAGGG SITTITGIGGG

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TOCOCOGGAGT GTTCAGTCTT GACCCTAGTC ACTGATTTTT TCTAGTTGTT AATAGAGTGG

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COTOMOTICAC CTCTCTATAG TOGGCOTOGC COMOGCCGGG GTGACCCTGC CGAAGCCTCC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 209:

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GAAAGAGGCY TITTICTCACA GCCATTATAT TAAATAGTAG GTCGATTCAC ATCYTCGTGC CCTTGTCACC ATCAGGCCTT TCTGGCTCCT GATAGGGTGG AGCAAAAGTG GAAAGGAAAA GIOSCOTOGG AAGGIGICCA CAGIGAGCCC IGIGIGCAGG ACTGICCACN ACGGITICACA

TOOTOGOOAC COTOCOOTOT GOOTOAGTGA CATGTAGATG ACTGAOTGCO AATAOTTGTO ACCATTCCCT GGAAGCAGCT ACCTAGGGGA AACAAGATOT AGTGCTATTG CCGATAACAA

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GTAAGATTTT CCACACTACA GCTGGGTGTT TCTCTTTTCT AAAGTGAGGC CAGTGTTATI

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TOCCAMOACC ATGCGCGGTG CTTAGGAAAT GAAAGAAGTC CCGGGTCTGT CTCTCTCACT TGATTGTGGT CTAATTTCCA ACCIGCTCTG TITTCTGTGA CATCITGGAG GEGGAGCTAG GGGGCCCAGA GGCCGCCTTT TGAAATGTTT GCCTGTCTGA ACTGTGAAGA CACTTGGGAG

CTCOCTCTCA MICOGOGAGG GAAAGAATOG CTTTOGTOGC TITTOTTCACA CAGCTGATOG

600 540 480 420

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(2) INFORMATION FOR SEQ ID NO: 209:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 932 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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AAAACTAAAA AAAATTAAAA AAAACTGGAG GGGGGCC

720 660

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GETCAGTGGE TGAACAGCAT TECCACAGCE TGEAAGTGTG TGTGTGTGTG AAAGAGAGAG

360 300 240

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CTGAGITIAGA ANTICACAAG TICTCCAGGT GATCTCATAC ATGCTAAAGT TIGAGAACCA

CATCACCATT ATCTOGNAAC ATGCAGTAAA TGCAGATTNT TCATCTTCTC CCCAGACCTC

GTTTCCCTGT TAACAAGAAA GTCAGAGGTC AGTTGATCAG ACATTAGATT ATTTATTGCT TIGAGIAAAG ITAAIGCATT AAGAAGAGAI TAGATAGGGA IGGIGGCGTA ICTICCTACA

660 8 540 480 420 360 300 240 180 120

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GARAGOGOGA CTCAGARGAA AGATCCTTGA CATTGCCWAA CATGCTGGGC TTGTCCAACA

180

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AMMACTOTO GACTITICOCA TATAAGGGOT GIGGITOTOT GIGGIOCCOT GGATIMGAGG

ATTAACTAGT GCAGTTGACT AATTCTCTTT ACCTTTATCA TTTARGGTGA RGCATTGCAC

120

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GAAAGITACA ATTOGAAAGT TICCIGCCAG CTICGGGAAT GACACTGCAA AGCTGATGCC

CCCAGCGCTG ATATGACAGT AATCCTCAGA GGCAGAGCCC AGCACAAAAC AGCAATGCTA

AGAAACTGCC AGROTAATTC TCCTCATTAC TGCTCTACCC ACCCACTTTC AGCTCCCCAA

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GITIGCITITICA GITOCCITITIA TITIGATICCI GGAGAGAGCA GACTICGCACS AACATICAAC

TACTICTAGG ATTATAAGGA ATTAACATIG AGAIGACATT TCCATTIGAG AAGGAAAATA

CCTCATTTTA GATGGGCCNC ANTATTTAAG ATGGACTGRG GMCCCCARAG ACTGACCCTT AANCCACTOC ANTITAAACC CCCTCCCCTC CAAGAAAGTT CACAACCGGC CATGGATGAC

CAGTGATGCG GCTCATCGAG AARCGGGCTT TCCMAGGACA AGTACTTTAT GATAGGTGGG

ATGCTGCTGA CCTGTGTGGT CATGTTCCTC GTGGTGCAGT ACCTGACATG AGCCAGCCAC

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SEQUENCE DESCRIPTION: SEQ ID NO: 207:

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(2) INFORMATION FOR SEQ ID NO: 207:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1166 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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	437		438
	GOTICOLAR ACCANTINCE ESCINEERS ENCYTORIS CONTINEED		GIGCITITICC TCGGTGGGAC AGTIGCTGGC CCTCTTAATT TTGGTGTATG TGCTTCCAAG
	מכרישורה פרוימוים שמנישורה פרוימוים	1.	TAICTAAACC TCCAGTCTGA TCTGTATATG CTATCCTAAC TGTTAATTGT ATTAITGATT.
٧٦	GCTCACCTTC AAGTCTATG CAGCACCAAA AAAGGACTCA CCTCCCAAAA ATTCCOTGAA 5	. 180	5 AIGTIGATIA ICTIGCTIGA AGGITCARAC TITICAATIT GAIAGAARA AAGITITITI
	GOTTGATGAG CITTCACTCT ACTCAGTTCC TGAGGOTCAA TCGAAGTATG TGGAGGAGGC	C 240	v
	AAGGAGCCAG CITGAAGAAA GCAICTCACA GCTCCGACAC TAITGGGAGC CAIACACAAC	. 300	
10	CTGGTGTCAG GAAACGTACT CCCAAACTAA GCCCAAGATG CAAAGTTTCG TTCAATGGGG	360	01
	OTTAGACAGC TATGACTATC TOCAAANGC ACCTOCTGGA TITTITTOOGA GACTTGGTGT	T 420	(2) INFORMATION FOR SEQ ID NO: 211:
7	TATTOGITIT GCTGGCCTTA TTGGACTCCT TTTGGCTAGA GGTTCAAAAA TAAAGAAGCT	480	(i) SEQUENCE CHARACTERISTICS: (A) LENTTH: GQ hase naire
?	AGTOTATICGS CCTGGTTTCA TGGGATTAGC TGCCTCCCTC TATTATCCAC AACAAGCCAT	240	(E) (C)
	COTOTITICC CAGGICAGIG GOGAGAGATT ATATGACIOG GOTITIACGAG GAIATAIACT	r 600	(D) TOPOLOGY: linear
20	CATAGAAGAT TTGTGGAAGG AGAACTTTCA AAAGCCAGGA AATGTGAAGA AITCACCTGG	099	20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 211:
	AACTAAGTAG AAAACTYCAT GITCTGCCAT CTTAATCAGT TATRGGTAAA CATTGGAAAC	2 720	GARACTIGACA TIGITIAAACA CACTAAAACA GAAGTACTTA CCTCTTGAAG ATTTAATATA
2,0	TCCATAGAAT AAATCAGTAT TTCTACAGAA AAATGGCATA GAAGTCAGTA TTGAATGTAT	780	TAATGGTTGA CATGATACAT GTACATGAAT GGAATGACCA GATGCTTATG GTCTACATTT
3	TAMATICOCT TICTICITICA GRAMAMOTA GACCAGACCT CTGTTATICIT CTGTGAMATIC	840	TCCTITATCC TGTIAGIAIT ACCTICCTIA ATCTITIGITC CITAACATGC TAAATICCTC
	ATCCTACAAG CAAACTAACC TGGAATCCCT TCACCTAGAG ATAATGTACA AGCCTTAGAA	006	TTCAGNETT ATTTTCTAGT GACAGAANGC TAACATTNCT TACACCCTGG CAGAAGGGAG
30	CTCCTCATTC TCATGTTGCT ATTTATGTAC CT	932 3	30 agadaistist titigggstgg staactadat tittgastga aatatcataa gatgagaatg
			GALAGAGGA GACACALAGA GITATRACAA AAAAACAATG GITITITIAG CCAITITGACT
36		Š	GOCTOTITINA ATRACTORCA AGACATICAC GITHUACATO ACTITITAGIG AAATAAAATG
ડે	HNT (7)	ń	TOCCATACTA GTATGTGCTT CAAAAGGGCA AATGTGCTTT AGTGCCCTAA GGCTAAATTT
	(i) SEQUENCE CHARACTERISTICS: (A) LENCTH: 661 base pairs		TOGICATITIG ACATCAGAGA TGTTGTAAGT ATTGCACTTA ATACGCACCT ATTTCTCAAT
40	(b) TYPE: nucleic acid (c) STRANDEDNESS: double (d) TOPOLOGY: linear		AGIGNIAITI TITIGGCIAG CAITINCIIT ACCACIAACC TIGITGGAIA GC
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 210:		
45	GICATICITI AARAAAGC TITICCTOTIT AAAGCITITIC AAAGAAGAG ACCACTIGA	. 60 45	(2) INFORMATION FOR SEQ ID NO: 212:
	AGATTCCCCC TAGGGTTGAT ATGTGTCTAA ITCATTITAT AAAARTAIT CITGTCTTCA	120	(i) SEQUENCE CHARACTERISTICS: (A) IDMITH: 918 hass naire
50	TITIPAAGCT TIGGCINTAL AGICAGAAT GICCIAAATA ACAAACIAIT FIGIATITAA	. 180 50	(a) (c)
	TTTAGGGAAG ACTAAAGGGA AGAAAATGA AAACTCAGTC TTTATGTAAG CTCCAAGGAT	240	(D) TOPOLOGY: Linear
	ATTAGGOCIT AAAGOOCITT TCTAGITTIA IGAGAAITTO TACTACIGAT TITITAIAIA	300	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 212:
55	TOCHOTITITI GAGARGACA GAICTCTGGG GAAARTOTTG AGTTACAATG GCAITTCACT	360 55	TOGAGTICOCT TECCACCTICA ATGAATCCTA TOTCTCCCCT GCAGGTICGTF GOTTTTCAAT
	GIGATECETE TEMAGETEMS ATCAGITETA TRACCEDATG ACAMENTOT TETTINGOTT	420	OTICHISCIA APPTIFIFICC TAPTIGACICT TGGGAGTITIN CPTIGITIGC TCCTGTGTT
09	. ACTOTCCTGT GAAATGTCAG CTCAAGTTTC CCAGAAGTCG TGTGTTTATG ATGAGTCAGA	09	GCCCAGCTIT AATAAAACCA GGCGCAAACA AAAACCATAG CATICTGAAA CAATAGGGGG

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TIGITICITG TOGCGAAATG TOGCTTTCAA ATTAAAATGM CCTTTTCTTC TTKGAAACT

360 8 240

CCCACATTOG ACCCAGTATO TCACTTTAAT OGACTTCAAG AAAAAATCTG AATOGGAAAA

IGACACTAGG AATGTATACT CCACACATTT TATOCCATAT AATGGTGTGT TITCTTAATI

PCT/US98/04493

WO 98/39448

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3 6 ઝ 25 20 55 8 30 8 5 5 GGTGTCCTTT CCCTGTCAAA CTCATTAAAA ATTCCTTT GCTGAAACCA AACCAGGCTT TTAAAAAACCT GTGTAGAAGA AAACCAAAAA ATCCTGTGTG CATTITICATT GAAGAGICIG AIGACTIGCT AGCGITTIAT CATTOTGACC ATOGTGATGC CTCATTTGCA TGATATGTAC CTTGTGTTTA ATGTGAAATA TECTOGRACE TRATTITISCT AMMIGICIST TRICITISTICA TRIGITISTAS CICACAGCAC TTATGACAGG CACACAGAAA CCATAGCATG GYCTOGCTTT CAGAAAANIGC CTCTCATCTT GCAGGAGGAA GGGCATGAGC CGAGCTTGAG GAATCCGTGY TCCAAACTCT ACACTCAAGG CGAGACACCC GAGGAGTCCG TTCCTCCCTG GTTACGTGGA CTGTGGAGCT GGTCTCTTGT AGCCTGCCGG GAGAGTGGTG GCATCTRARA GOCTGGTCGT GGACTGTGGT TGGGGGGAGGT AAGTACATGC CTCCACATAA TGCGGTGCTG TCCATCTCGG CAAATACTGG CCAAGTCCCT AACCAGGAGA TACAAAGAAG TCTCAGTAGT AATCTTGTTC ATGTGCTTTT ACAGCCAGCT TOCCCAAGAT GCCCACACAG TCGGAGGTOG ATAACGTOTT TGACACAGGC TTGCGGGACG TOGANCOCOO GEOCTECTEE AGETTECONG TECNOCENOE CTOGGEOGEO GOCGEOCECE BOURDETT TIMACCOTOT OCCCCCTCTC CTOTOCCKGC GTODOCATCC CCCBBGCAG (2) INFORMATION FOR SEQ ID NO: 213: ACATTIMAGR ATGINITAGI TACAGAAATI ATATGICIGI GINIGIGICI CTACICAATI TTTGTTTKGA CTKGTATAAT TAAGGGTTTG GAAAGATTCA TAATTMTGAG AGAGGTTTGC AGAAGAAGTC TCCCTGGCCA TTCTCAGACG AGTGCATCCC ATGGGAAGTG TGGACGGTCA RTOCMITGCG CAACTOTGGT GGCGATGGGC TGGGGCAGAT GTCCTTGGAG TTCTACCAGA GOCTICAGOOC COTOCOGAGO TTOAAGCOTA CCTOCOGAGO TCOCACCAGO GOCOTGAGGA FIGAGAAACT CIGCGAGAAG AICAICAACA ICOIGGAGGI GAIGAAICGG CAIGAGIACI AGGIOCAIGI GGIAGCCCIG GCCACGGAGC AGGAGCGGCA GAICIGCCGG GAGAAGGIGG E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 213: (B) TYPE: nucleic acid (D) TOPOLOGY: linear (C) STRANDEDNESS: double (A) LENGTH: 1079 base pairs TITTICIGIA AGCICAAIGI 900 660 60 540 480 420 360 300 180 120 938 840 780 720 660 600 540 480 420 8

25 20 5 5 S TIGGTTTTGT GGTTGCCAGC CTCAGGTCAT CCTTTTAATC TTTGCTGACG GTTCAGTCCT (2) INFORMATION FOR SEQ ID NO: 214: AOCCACCACT GGGATGGGGA ATAAAGTTGA GAACATGAGT TTGGGCTGAA AAAAAAAAA SCOTOTACTS TOTOTOCATA SCOOTSSTOS SSTOCOCOOTT OTTTOTOCAC TOTACAGAAS GCTCTGGAAC CTGCTCTGGG TCATTGGTGA GACTTGGAAG GGGCAGCCCC CGCTGGCTTC GGAGCICCI TGATGGCICC CAGACCITGG CITTIGGGAA TIGCACITIT GGGCCTTIGG CCACCACCAT GCGCAGGCTC ATCAAAGACA CCCTTGCCCT CTGAGCGTCG CTGGATCTCT TOCAGECETA CETOTACAAG ATETECETTEE AGATEACTGA TGECETOGGE ACCTEAGTEA (1) SEQUENCE CHARACTERISTICS: (C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 3791 base pairs

> 1079 1020

960

900 840 780 720

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 214:

ઝ ઝ 3 8 8 55 8 QUANAMADAA GCCGGCGGGG GCGGCGTTGG GGGCCCTGGG GCCAMGMGCG CATCMGGGCC AAATOTOACT GAGGAAACAC CTGAAGGTGA AGAACATCAT CCAGTOOCAG ACACTGAAAA TODANGTONT AGAICTICTT TITICACATTA CAGTOGCCTG AAGCACGAGG ACAAACOTGG TODANTOOOC CONCONGNIO GATTTOATTC TCOTOOCAAA COTGAATTTO ATAGGCATAG TATTGACCGA CCTATTCGAG GICGIGGIGG TCTTGGAAGA GGICGAGGG GCCGIGGACG ATTCGAMAG CCACTTGAG AMMGGGTGA AGGAGGCGAA TTTTCAGTTG ATAGACCGAT TCAGGGTGAA GOGAAAATAA TTGATAGAAG ACCAGAAAGG CGACCACCTC GTGAACGAAG CCCGTGGCGC TTTAAGAAAG AAGGAATAAG ACGAGTTGGA AGAAGACCTG ATCAACAACT COCAMGAACC COCTOCCCC CAGOOTTOOC OTOOTTOACA AGAAAGAGGA GACOCAGCCG GOGGOCCAGA CCAACTOCAA CGOGGAGGO AAACAGOTGO GCAAGGAGTO CCAGAAAAGAC CGACCAGTTA TITIGACGACG AATCOGACCC CTTCGAGGTG CTGAAGGCAG CAGAGAAACAA AGCCACCATC ATOCCTOGGC ACTTACAGGA AGGCTTCGGC TGCGTGGTCA CCAACCGATT TEAMOCAGOS GETETTOSOST COSCOCOSOS COSTOSCAMIC COTOGAGGAA COCOCOSOS OTOGRAGOCT ATTICARARTA AGGRECOGGE RARAGTAGRA TITRATRITCE GRARACCIAR TAAGGAGAAT GAAGTTGAAG AGGTAAAAGA GGAGGGTCCA AAAGAGATGA CTTTGGATGA AGGTAGCOGA TCTCACAACT GOOGAACTGT CAAAGACGAA TTAACTGACT TOGATCAATC 540 360 300 240 180 120 660 600 480 420 960 900 840 780 720

PCT/US98/04493

TGANGGTGCT GANGGGGGG ATTTCTTCTT CATABATCAA AGAGTGAAGA GGCTCANGCT GAAGATTCGG TTATGGACCA TCATTTCCGG AAGCCAAA ATGANTAAC GTCTCAACT GAAGATCAATT TTGAAGACCT TGGCCGCCCA GGAGGAGGG GCAGGGGAGG

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1200 1260 1320 1380

ACGAGGTGGA CGTGGCCGTG GTGGCCGCCC AAACCGTGGC AGCAGGACCG ACAAGTCAAG

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TOCTTOTISCT CCTGATOTISC ATGACCCAGA GGCATTOCCA GCTOTISCTT AACTGAATOC CATAAGACAA CCCTGGTTOC TTTGTGAACC CTTCTGTTCA AAGCTTTTGC ATGCTTTAAGS

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1500 1560 1680 1680 1740 1860 1920 1920

AGACTGAATT TTATCTGTTT TAAAAATGAA CTTCTCCCGC TACACAGAAG TAACAAATAT

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GATTATIGCAT TCTTCATGAA TACTTTTGTA TTGCTGCTTG CAAATATGCA TTTCCAAACT

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GGTAGTCAGT TITGTATTTA GAAATGTATT GGTAGCAGGG ATGTTTTCAT AATTTTCAGA

ANTIMAGENT TREGMETTC CCCCANTRC AMCTEGITT TRANTINGS ACATACTGGT
TITAMINCT GCITICCATA TECACAGE GICAACTGGG ACATOTINA CITICATITG
TCAMITITA TECTOTOGGG ANTACTACT AINTGINITT TRACTINGIT TITATATITTT
CATTITIGGG GAAAAATCTT TITICACTTC TCATGATAGC TGTTATATAT ATATGCTAAA
TCTTTATATATA CAGAAATATC AGTACTTGAA CAAATTCAAA GCACATTTGG TTTATTAACC
CTTGCTCCTT GCATGGCTCA TRAGTTCAA ATTATAACTG ATTACATTT TCAGCTATAT

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TGAAATATAG GTGTGAACAG TGTGTACCAG TTTAAAGCTT TCACTTCATT TGTGTTTTT

ATTICCADACG ACTAAGADAT TAADADAAAA AAGACTOTICA TTICATACCAT TICACCIDAA

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	ATTIAGCIGA TIGGITCICA CATATACTIC TAAAAGAAAC TITTIAIGITA TAAGAGTIAC	2820
	ITITIGGATA AGAITTAITA AICTCAGITA CCTACTAITC TGACATTITA GGAAGGAGGI	2880
\$	AATTGTTTTT AATGATGGAT AAACTTGTGC TGGTGTTTTG GATCTTATGA TGCTGAGCAT	2940
	STICTICACT GOIGCTAAIG ICTAATATAA TITTATATIT ACACACATAC GIGCTACCCA	3000
	GAGAITAAIT TAGICCAIAT GAACTAITGA CCCAITGITC AITGAGACAG CAACAIAÓGC	3060
01	ACTECTIAAT CAGTOTOTT AGACTITICA AGTATCTAAC TCATTICCAA ACATGTACCA	3120
	TOTITIATAA ACCICITGAT TICCAGCAAC ATACTATAGA AAACACCIGC TACTCAAAAC	3180
15	ACAACTICIC AGIGICATICC ATTGCTGTCG TGAGAGACAA CATAGCAATA TCTGGTATGT	3240
	TECHACTITI CHAGNINGCC TGANCTINAA AMGTIGGIGC ATINGITGIN TCTGATGGAT	3300
UC	ATAMETTICS CTCCTAGTIC ACTITICIOTIC ANGAGCTANA ACTICICANCE TANCITICIC	3360
07	TIATIGGIGG GIAATAACIG AAAATAAAGA TITATITICA TOCICACITC TIAAAAGICA	3420
	THAMAGAAT CAAATAGGRT CATGTTTATT GTCATGTGTT TCCTGGRTTC TGACCTGTGT	3480
25	GCACACCCCT GTGTGTTTAT ANTTITTAAA TTGAATTTTA TATGGGGTTT TTAITTTGCTA	3540
	AMANCCAGGC TGTTGAATCA CATTTGGGAA GGGTACTTAT CTTAATGACT AATGACTTAA,	3600
02	TIGGGAAGT TGAATTCTIG TAAAATACAA AATCCAAGGA CTTCTTGGGA TTTAATCTAA	3660
	TIGICACTIC NITAGGCAGA INCACTITIT IGGATAATIGG AAAGTTAAGC ATACCGAATG	3720
	CTACTTTTGG TTGACAAACG GGCCTAATAG TCCGGGGGGA AATCCCTAAC NGGTAAGGNT	3780
35	CCCAAGIATO G	3791
40	(2) INFORMATION FOR SEQ ID NO: 215:	
45	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1314 base pairs (B) TYPE: nucleic acid (C) STRANBERNESS: double (D) TOPOLOGY: linear	
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 215:	
50	CAGIGGICOC TECTGETEGG GCGCTGGG CECCGGGCGT CGCCATGACT AGTCAACTGG	09
	ACAICTICGT GGGGAACAGA CCCTTAICGA CGAGGACGTG TATCGCCTCT GGCTCGATGG	120
\$5	TINCTICATION ACCONCICOS TOSCOCTICOS GOTISCICOS GONATOCTICO AGCAGACTICO	180
•	COCCACOGOA GOGOTACTNC AGAGGACAC CATGGACCAT TACCOCACCT TCCACATGCT	240
	CGAGGGGCTG CTGCATGGGC CGCCCAAGCT ACTGCACCAG YTCATCTTCC AGATTCCGCC	300
09	CTCCCGGCAG GCACTACTCA TCGAGAGGTA CTATGCCTTT PATGAGGCCT TTGTTCGGGA	360

2040

THACTITITA AARGETIGAG TITOCCAITT TAAAAICHAA ACHAGACAIC TITAATIGGIG
AAAGTIGITT AAACHACHA TIGTIGGIAG GCACARGGIG TCAAGIGAAG TAGTITIANA

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GTAATATTA AACTOTTCTG TAAAATAAGT GTCTGGCCAT TTGGTATGAT TTCTGTGTGT GAAAGGTCCC AAATCAAAA TGGTACATCC ATAATCAGC ACCATTTAAC CCTTCCTTGT TCTAAAACAA AAACCAAGG GGGCTGGTTG GTAGGGTGG GTGGGGGAGT ATTTTAATTT TTGGAATTTG GGAAGCGGAC AGCTTTACTT TGTAAGGTTG GAACAGCAGC ACTATTAATTT TTGGAATTTG GGAAGCGTT TACTGTTTCT AAATTTCCTA GATTGCTATT ATTTGGTTGT AAGTTGAATA TGGATTCCTA TAATGGGAGC ATCACCACTT ATTAAAACA CACAAGAATTG ATGAATTAAA AAAGTTTTCT AGGATTGTCT GTTTATTCTGC CACATTTTATT GATAAAAGAGT GAAGGAATTT TTAAAAAATT TTTAAGAATT GTTTGTCGACG TCATTTTTAG AAACATTTCTA

GOTATGGGTT TITTCTCCCC CTTCACCAGG GTGGGTGGAA TAAGTTGATT TGGCCAATGT

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GGTGCTGGGC AAGAAGCTGT CCAAAGGCAC CAAGAAAGAC CTGGATGACA TCAGCACCAA

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STACSGATOT TAASGACTST SCCATTOOTS COSCISCIAS SIGSCISCAT TGATGACACO

COCCUTTUTO TOGGGCACAA CATGCTGCTG GTCAGTGAAG AGCCCAAGGT CAAGGAGATG RICTICCCCA ACCAGGICCT GAAGCCCTIC CIGGAGGAIT CCAAGIACCA AAAICIGCIG

ANCONCAGOA GGCAGGGCTT TANCAACTAC TOCAAGCTOO CCAGCOTGCO COTGGTGCAG

AGGGAAGAAA AAACIGCAGT AICIGAGCIT COGTGACTIT GCCTICIGCG CIGAGCICAI GGTGGTAGAG GAMATGCGGG GCTCCCTGGT GGACAATATT CAGCAACACT TCCTCCTCTC AACAGGCATC ACCCTCAAGA GCTGCCGOAG ACAGTTTGAC AACITTAAAC GOGTCTTCAA TGACCGGTTG GCCAGGGACT ATOCAGCCAT CGTCTTCTTT GCTAACAACC GCTTTGAGAC

S

AGACAAGGAA TITICICCAGG ACTIGAAGGA GCICAAGGIG CIAGIGGCIG ACAAGGACCI GATCCAAAAC TOGACCCTTG GAGCCGTCGA CTCACAGATG GATGACATGG ACATGGACTT

> 660 600

> > 5

GOGGAGCTTG TAGGAGGCCT CACCTGCCTC ACAGCCCAGA CCCACTCCCT GCTCCAGCAC

660

600 540 480

720

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TOTOGRACOTO CACARGRAGO TOGTOTOCAO TOCTOTOCO GGRARAGOTOG GOGTOTTOTO TGAGATGGAA GCCAACTTCA AGAACCTGTC CCGGGGGCTG GTGAACGTGG CCGCCAAGCT

GACCCACAAT AAAGATGTCA GAGACCTGTT TGTGGACCTC GTGGAGAAGT TTGTGGAACC

960 900

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CTOCCOCTCC GACCACTOGC CACTCAGCGA COTOCCGTTC TICCTGAATC AGTATTCAGC 1080 1020

GTETGTECAC TECETEGATG GETTEEGACA COAGGETTET GOGACEGETA CATGGGCACE

25

AAGTIGCTCT GAGTTTGGAG ACTOGTCCTC GCTCCGGGGA GCAAGTGGGG GCCTCCTGCG CCTGTATCAT GACTGAGGTG CCTCCCAACG CTCCGCCCAC 1140

TOTOCCTOTO TCTGTCTCTO AGCACCTGGT GTCCGTGTAC AAGGATGGAT 1260 1200

GIGINCNOIG GCICCTIGGG AACIGAGACA TAICICAGGG AAIGGIGICI GIGCICAGCC 1320

30

CAGAACAGOG ACTTGAAACCA AGCCCTCTGC TCTGAAGACC GCGTCCTGAA TTTCTTCACT

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TGITAGTICT CCACCCICGA GGIGTACGCT GIGAAAAGII IGGGAGCACI GCIITATAAI AGAGOTTOCT CATCAGGITA COCAGAAGIG GGTCCCATCC ACCATCCAGG TGIGCTIGGA

1500 1440 1380 1320 1260 1200 1140 1080

1511

(2) INFORMATION FOR SEQ ID NO: 217:

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(1) SEQUENCE CHARACTERISTICS:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1511 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

2)

INFORMATION FOR SEQ ID NO: 216:

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CATCCACCAG AAGA

1334

(A) LENGTH: 642 base pairs

(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 217:

GACAMAGITC ACGTAGCAGG TCTAGGCAAA GACTGGGCAA TTGAGCAGAG GAGACGGACC AGGCCTTACT TITICCTCCCA CAAAGGAGTC GCAGCCACGC TAGCTCTGAC TTGCCACTGT ACCRYGAGSC GGRCCCCTTC ACCTTGGCTG GGCTGGTCCT GGTCCTTAGG

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WO 98/39448

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TOTTTAGCCA GOCCTGCGCA TANATACACT CTGCGTTATT GOCTGTGCTC TCCTCAATOG

TOTATION TO COOCCUATOS GAAGOCIAGAT COTGACACTO TROCOGACTO GTAGOCIAGOC CAGCCCCTCC AGCTGACCAC CCTGTTGGAC CAGTACATCA GAGAGCAACG CGAGAAGGAT

GACATOTOGA AGAACTIOGO GICOGOGAGI GIGITITOTCA CIIGGITITIC ACTAGIAAIG

900

20 ATATTOTCAG GTATAGOGCC ACTTOGAGAT GCAGAGGATT CCATTTCAGA TOTCAGTCAC

COGCTICGIC CITAGITITIC CCAACITOGG ACGIGATAGG AGCAAAGICT CICCATICIC

1020 960

CAGGICCAMG GCAGAGATCC TGAAAAAGATA GGGCTATTGT CCCCIGCCIC CTIGGICACT

GCCTCTTGCT GCACGGGCTC CTGAGCCACC CCCTTGGGGC ACAACCTGCC ACTGCCACAG TROCTCAACC AAGCAGTTOT GCTGAGAATG OCACCTOGTG AGAGCCTOCT GTGTGCCAGG

25

CTITICIOCIG AGIOCIGIAC AIGIAITAGI ICCITTIACIG CIGACCACAI IGIACCCAIT

TCACAGAGAA GGAGCAGAGA AATTAAGTOG CTTOCTCAAG GTCATGCAGT TAGTAAGTOG

30

180 120

TITIOTCAGG TIGICCTIOT TIGGATCCCT CAACTAGGTG ATAAGCACTG GAGGGGGATG

240

8

360 8 240 180

8 AGTGCAGAGO ACAAGCTTCT TATGCGACAC CAGCTGCGGA AACACAAGAT CCTGATGAAG

55

CTOCCATCTC CTCCCAGCCC CCCACAGGAG GAGATAGGCC TCATCAGGCT TCTCCGCCGG

GAGATAGCAG CAGTITITCCA GGACAACCGA ATGATAGCCG TCTGCCAGAA TGTGGCTCTG

CAGAAGCTGA TOGCTOTGAC TGAATATATC CCCCCGAAAC CAGCCATCCA CCCATCATGC ACTOTOCOCT ATOCCTOCAA GOCTOTTACO COCCACOGTO OTOTGATOCA CTITICAGOGO

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GIGGEGGGA IGCIGEGAGG GGGICICCIG CCCEAGGEGG GCCGGCIGEC TACCCICEAG

SEQUENCE DESCRIPTION: SEQ ID NO: 216:

	WO 98/39448	PCT/US98/04493 W	WO 98/39448
	445		446
			CHOCAGITAC CATAATITIC CATGITIGIG GAATIGATAT TGAAATAGCA GGGCTAAGGA
	ACCCOCCTIG GACGIGITIC TITAACCICA TCCATATAAT AGGCCGTGG GATGGTTGTA	300	ATTACTOSCA AGTITITACC TOTOSGRAT ACCTIAGOCT TATITIAATA TJTGTAATT
ν.	GAGGTAAAGC AGGATGATGG TOTTTTAAGA CCAGAGCTTG GGACCAGGGC TCCTACACCT	360 5	TATTIVAATG TICAIGAATG TIIGAAAGGA ACAAAATTAT CAGGGAIGGC TCTTIGGCA
ı.	ANITHETET CETOSTAGET GAACAAAGST CTAAATTAGE TTAACAAAAG AACAGGETGE	420	GGOTCTTAIT TICACCCICT TITCTGTAG AAAAAAGAAC AATGICTTAA TGIAITTITA
	CSTCAGCCAG AGITCTGAAG GCCATGCTTT CAGTTTGCCT TGTTGACAAT TGCTCTCCAG		AACTITITICS TATACTITICE AATTICEAATT TITAATAAAAG T
9	TTCCTATGAA AGCACAGAGC CTTAGGGGGC CTGGCCACAG AACACAACCA TCTTAGGCCT	540	
	GACCTOTIGNA CACCAGGGG TTGTOTOTOT GITCTOTITIC TCTCCTTTGCC GAACTITICTC	009	
2	AATAAACCCT ATTTCTTAIT TTANATTTAC GINGGIOCTG GG	942	(2) INFORMATION FOR SEQ ID NO: 219:
2		2	(i) SEQUENCE CHARACTERISTICS: (A) LENOTH: 1080 base pairs (B) TYPE: nucleic acid (C) STRANDENESS: double
20	INT (2)	20	
			(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 219:
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		TOTTIATOTS ACCTARABGA TACACACATS CACACACA TACATATCCA TTCATTCATT
25	(D) TOPOLOGY: Linear	. 67	CATICAAGIG GIGITICCAG IGICIGIGIG ICACIGITIA IGCAGITICC AFFICCCAGI
	(*1) SEQUENCE DESCRIPTION: SEQ ID NO: 218:		GAATTATGAG TGGAGGCAA CTTTTCTAAC CAGATTGTCT TTTCAGAACA AAGACCKGGG
30	GENCEACTG TICCAITITA INCIPATAGA TICCAITICTA GGOCCAGCC GICTCTIGAC	90	rattgaggaa gagtttggaa agagggaga gcaaggaaag agagctttaa attgaaaggt
	TGATGOTGTT CCCTTTAACC CTTGGCATGT ATAATAGAAT TTTGGTGAAT GAAAGAACCC	120	TAATTICCIA AGAGGAACCI GGGCIGAAIG ACTACAGIGI TAIACCCICC AATCITIGCA
	AAATAGGCCA GATAGTCCCC CCAGGCCCTG ATATCCATAA AAGGCTTGGG AATGCATTAT	180	GGTGGGCATG GAACACTGCT TGTATCACTC TGTGCACGGT ATAAATCCAT ATATCCACAA
35	GTAATTGTCC TTAGTCTTTT TGTTGTTTTA GAAAAAAA ACAAGATGGG CTCAGATGGA	35	AAACACACAT CCATCCATCA ACATATACAT GOTTTGGGAT GAGCAGGTCA ATAGTTTTGA
	TOCCTACOTA AAAATGOTIC CTACCTGTOT ACTCATAACT TITCTTTGAA TTGAGTAGTG	300	GAGGAGITI GITCCITITI ITITCICATI AFACICITAA AITGITGICA GITATCAAAC
. 6	AAAGGAAGGA GGAGGAAAGG AAATTAAATG TOCTTCTAGF ATTOTOTGA CTCAAGTCTG	360 40	AAACAAACAG AAAAATTGTT TGGGAAAAAC CTTGCATAGG CCTTTTCTAT CAAGTGCTTT
	ACATATGRGA TAATAACCTA TATTGAAATG CCAAGAATTG TATCTGAAAC AAGRGAACAG	420	AAAATHINGA CTAAATACAC ACATCCTGCC AGTTTTTTCT TACAGTGACA GTATCCTTAC
	TITGACACAT TIATCATGCC TTCATATTAC ATAITAACTG AAACCAATTA ATAAACATAT	480	CIGCCAITTA ATATTAGCCT CGPATTTITC TCACGFATAT TTACCTGTGA CTTGTATTTG
45	GAMATATICEA TTGCACAAGG CAAAGGCACC TAAACCTETT GTTTCTTTTT CTACATAGCA	540 45	TTRITTAAAC AGGAAAAAA ACATTCAAAA AAAGAAAAAT TAACTGTAGC GCTTCATTAT
	GAAAITGAIT TITITITAT TITITIAGGG GAACCTATAT AATTATGACC CAGTGAIGTC	009	ACIATIAIAT TATTATTATT ATTOTGACAT TITGGAATAC TOTGAAGITT TATCTCTTGC
8	TTTTGGTGAC TTAAGCTTAT GAATTCAGGT TACAATTGAG TTGATTCTAG ATGGTTACTA	909	ATMINCTITIA TACGGAAGTA TTACGCCTTA AAAATACGAA AATAAATITIT ACAAGGTTTC
	CCTTGANAAG GATGTTRGTIG CCTTATGTGA CACGAGCCAG AGCCTGCTGG GAATAAACAA	720	TOTITIGICI GGAAGAGIAA TIGANGINGC TAAGAATGAT GITNGTITIT TIGGGGTITIT
	AGCAGATTCA TOCCAACACC AACTCGTAGC TTTAGTGGCA GATGGGAGTG GTCACAGACT	780	TOTTOTITT TITTIAAAIG TIACCAGCAC TITTITIGIA AGTITCACTT TCCGAGGIAT
22	CCCAAAAIGT GGGGCTTTGG ATTTCCACAC CATCCCAGGT GTGTGAUC TTCCTCTTTC	. 55	TSTACMOTT CACACTISTIT STGMSSTITS MAINTGMSS MAINATIAMA MAMAMAMA
	ACACTOTTGA TGATAAITTG AAAATGRTGA AATCACCTCT GAATTTGOCT ATAGCATGAG	006	AAACONCOSO GOGGOCCOCO TECECATIVOSN ECEAAAGOGGO COGTIACOCO GICAAAGGCCG
9	CACATTCTTA TGACACATA ACAAATAGTT CATAATGTGA ATATTAGAAA CTOTTACAGC	09 096	

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447

INFORMATION FOR SEQ ID NO: 220:

2)

Ê SEQUENCE CHARACTERISTICS LENGTH: 1258 base pairs

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 220:

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TECCTATITE ATTICACITY ATCCCTATTA ATTITITACA GIGAAATITY ATTAAAGTAT TGAATTGAGG GCTTAAAGAT AAACATATGG GRTTGGAGTT GTOTGTCCAT AGGGTTTCAC 120

180

5 5 AMARCITICAT CCACATCACA CCCIGITITAT TITICCTITAAA CAICTIGGAA GCCTAAGCTT AGAGCAACAT TOTOTTAITA AAGCATAGTT TATTTCACTA GAAAAAAATTT AATATCAAGG CTGAGAATCA TGTGGCAAGT GTGATGGGCA GTAAAATACC AGAGAAGATG TTTAGTAGCA CACAMINIAI GAAAIAGIAC CCICIAAAAA AGAGAAAAA AAAAICAGGC GGICAAACIT ACTATTACAT ACTICATTAC TAGGAAGTIC TITITIAAAAT GACACITAAA ACAATCACIG

> 180 120

SEQUENCE DESCRIPTION: SEQ ID NO: 221:

(A) LENGTH: 1693 base pairs
(B) TYPE: nucleic acid
(C) STRANDEINESS: double
(D) TOPOLOGY: linear

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WO 98/39448

INFORMATION FOR SEQ ID NO: 221: SEQUENCE CHARACTERISTICS:

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GITTICICITG

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IGGCTACTAG AAGTOTCCCA GAAGTCACTG TATTTTTGAA ACTTCTAACG TCATAATTAA

ACATCTGATT CTGGAATGCA GAAGGAGGG TCTGGGCATC TGTGGATTTT

TCTTGGGCAT CAAGANTAGT TCCAATTTTT TGGGCCGGG CAGGGTGG

1258 1200 45

TOTGAGOTTA TITTICCATT TGATATICAT TGATATCATG ACTICCAATT GAGAGGAAAA TITTICICAGA AGGATKAGAA ATCAGCACCT OCCITITIAGA GATCATAATT CTCACCTACT GTGCCAGTTA GTGCCTTCGG TGTAAGATCT TCTCATCAGC CCTCAATTTG TGATCCGGAA AACCCAKGTG CATTICIGCA TCICCIGGAT TAGCCITISA CAIGITGCIG RCICACATTA AATTOTOTO TGACTACTOT COAGTAAGGA GGCCCATTOT CACTTAGAAA AGACACCTOG

IGAGATCAAA TGTCATTTCC CAAATTTCTT GTAGGCCGTT GTTTCAGATT CTTTCTGTCT

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ACTOCCCTAC TCAAGACGCC CTAAGACTGG TAGAAATTAA AAGGATTTCA AAAACTTTCT TTATOCAATO GTTTOTTOAG ATACOGACIT GATGOTGCTG TTTAATCAGT TTGCTTCCAA TATIGAGGAA AGGTATICIT CYATACAACT IGITTIAACC TITGAGAACA TIGACAGAAA GAGAATTCAG GATAGITITIG TITAAAITICT TOCAGAITAC AIGITITITAC AGIGOCCIGC TTATTTAATT TITTÄGGTAA TGCCTATCTC TTOGTCTATT AAGGAAAGAA GCAATCAGTA TICAANIACC CAAAITOIGA TAGCAIAAAI AAAGIATIIA TITTAIGCCI CAGIATAITA GCTGGCACCG TTGCTTCCTC TTTGGGAAGA GGAAAGGGTG TGTGAACATG GCTAACAATC GCATATTGCA GAACAGCTCT GAGAGCAACA GITTCCCCATT AACTCTTTCT GACCAATAGI GAAAAAAAT CATTTTATCC GICTTTTAAG TATAIGTTTA AAATAATAAT TTAIGIGICT

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GAGGICTITT AAGICTCTIC CCTTIGGTIG CCCACCIGAC MATTITIATIA AGTACATITIG

AAGTECTECA GETOGTECTO CIGCTAGTAG TGTTTGGYTT ATTTTCCATE CCAGTICTOG AATTICCACAG AAGATIATICAT GTCTTTGTCC ACCCAAGGGG ACTACATICAA TCTACAAACC GTCAAGGAGA ACAAAAATAT TAATGAGGCT ATGAGAGTCC TCATTGAAAA GATGATGAGA COGGASCAGA TIGACCOGIT CAGTAAAGAG AACGGTITCA CAGGITOGAC AGAAACAICA GAGCCGGTGC CCTGCCTGCT CTTGGCCAAC AAGTGTGATC TGTCCCCTTG GGCAGTGAGC TTCAGCAACA GCCAGAGGTG GAAACAGGAC CTAGACAGCA AGCTCACACT ACCCAATGGA TIGIATTATO GGGAIGCOIC IGCOIGIGIT AFTAIGITIG ACGITACCAA IGCOACIACO ATAGTOCOGO TICAGOTOTO GOATATIOCA GOOCAGOAGO GOTTCACCTO TATGACACOA ANTOTACATA TATTITICAGI GGATTITIGCI CIGAAGGITIC ICCAGIGGIC IGACIACGAG

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8 ATAGAAGAG CCATATATAT TOCCICCTTA TCCTTGAGAT TTCACTACCT TTATGTTAAA TITICCAGIGI TGAATCICAC ACACTGTACT TIGAAAATTT CCTTCCATCC TGAATAACGA

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GTATTTATTO TOCTOTOTOT GGTCCTAAGT GGAGCCAATT AAACAAGTTT CATATGTATT

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TATTCCCTTT AAGAATCTTT ATGTATCAGT GTGAAGATCC TAGCGAACCT ATGCTCAGAT GAAAGITTIGA CCAAATITTOT TITTITTOTIG TIGITGITGI TITGAATITIG AAATCATICI

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	449		450
	acticiciat aatictitaaa alcichgaa gaataaaag iggaititaak ittaaaaaa	1680	(2) INFORMATION FOR SEQ ID NO: 223:
v	padadadada dad	1693	(i) SEQUENCE CHARACTERISTICS:
. و			TYPE: nucleic acid STRANDEDNESS: doub TOPOLOGY: linear
0	(2) INFORMATION FOR SEQ ID NO: 222:	01	
2	(1) SEQUENCE CHARACTERISTICS:	1	
			TCAGGAAGT GGCAGGAAAG GCTTGGAACA GCTGCCGGAG TGACGGAGCG GCGGCCCCGC
2	(C) STRANDELNESS: double	·	CCGGTTGCGC TGGAGGTCGA AGCTTCCAGG TAGCGGCCCG CAGAGCCTGA CCCAGGCTCT
<u>C</u>	(D) TOPOLOGY: linear	2	GRACATYCTIG ACCCCAAGTC CCCCACACTC AGTOCAGTGA TGAGTOCGGA AGTGAAGGTG
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 222:		ACAGGGCAGA ACCAGGAGCA ATTTCTGCTC CTAGGCAAGT CGGCCAAGGG GGCAGGGCTG
20	ACCCOTCOCT CCACCCACGC CTCCCCCACN TCCCCTCGTG GGGAAGGGAG AAGCATTTGT	60 20	GCCACACTICA TICCATCAGGG GATTGAAGGC COTGGGGGGT AGGACTIGGTG
3	АААССССВЕА ВСВАБЕТТСТ ВСТТАСССВА ВОССОСТВСТ ВТВОЗВАВАЕ СОССОВЯЕТСЯ	120	CAPARICYCE ARCHITERAR CARCONERS REPORTED FOR A PERCANDING CONTRIBUTION
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ć	TOTTCCAATG AATTCACTCA GOTTTCTCTT TGAGGOTCAG AGAATTGCTG ATAATCATAC	360	מוששתמומו וההרשושות שמומוותרות השתמורוותה הרומתמישו מומרמת-שמה
ફ	TOCAAAAGAA CTGGGAATTGG AGGAAGAAGA TGTGATTGAA GTTTATCAGG AACAAACGGG	30	INGARIACCT INTRATIGAG GCINIOTATO CIGACNICCT ICONGCICC CINGARCAGO
	GOSTCAITCA ACAGITIAGA TAITCITITI ATTITITITC TITICCCICA ATCCITITIT	680	GCAACCAGCG GCTCGAGGTT GACTACAGCA TCGGGCGGGA CATCCAGCGC CAGGACCTCA
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40	TICATIGIGE TGATTITIGG TGATCAAGCE TCAGTCCCCT TCATATTACE CTCTCCTTT	660 40	COGCAGCCAC ATCTCAGGAC CCTGAGCAAC ACCTGACTGA GCTGAGGGAA CCAGCTCCTG
	TABABATTAC GTGTGCACAG AGAGGTCACC TTTTTCAGGA CATTGCATTT TCAGGCTTGT	720	GCACCAACCA GCGCCASCCA GCAAGAAAGC CTCAAAGGGC AAGGGGCTCC GAGGGAAGCC
9	GOTGATAAAT AAGATCGACC AATOCAAGTG TTCATAATGA CTTTCCAATT GOCCCTGATG	780	CARGATTING TECAAGTEGA ATTGAAAGRA CTGTCGTTTC CTCCCTGGGG ATQTGGGGTC
5	TICTAGGATG TGATTACTTC ACTCCTGGAC TOTGACTTTC AGTGGGAGAT GGAAGTTTT	640	CCAGCTGCCT GCTGCCTCT TAGGAGTCCT CAGAGAGCCT TCTGTGCCCC TGGCCAGCTG
	CAGAGAACTG AACTGTGGAA AAATGACCTT TCCTTAACTT GAAGCTACTT TTAAAATTTG	006	ATMATICCTAG GITCATGACC CITICACCTCC CCTAACCCCA AACATAGAIC ACACCTICTC
20	ACCOTETIGA CELLALAGALG AGSARTRICA COTTICAAGTE AAGRIGACHG RIAAGGTGAG	960	TAGGGAGGAG KCAAATGTAG GTCAATGTTT TGTTGGFACT TTCTGTTTTT TGTGACTTCA
	AGTAATGACT AACTCCAAAG ATGGCTTCAC TGAAGAAAAG GCATTTTAAG ATTTTTTAAA	1020	TO A LAW COMMAND TO A LAW COMMAND COMM
	AATCTIGICA GAAGAICCCA GAAAAGTICT AATTITCAIT AGCAAITAAT AAAGCIAIAC	1080	MONOCONTO DESCRIPTION DESCRIPT
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TCCACTIGCA GITCITTAAA GAATGCIGCT

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ACGIGGCCAT CIGGAGCIGG TGCTATAGGT GACCATCIGG TACATIGAGG

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CTGGCCCAAG

AGAGCATCCA GAAGGCAGTA GGACCTGGTT

TITCAGGIAC TGGGAGCCGG

CAGACTICIC GITAGCAGCI GGAAGACATI CCICCCACAC TITICCCTIC

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AAACCAGAGA CTCATGTTTC CAGGGGTCAG TCTGTCAGGC AGGAAGGACC CAGGATTTGA GOCACATGCC GAAGATACTC AAGAGCTCCC AAGATTTGCT TGAGGCTAGC CCAGTGAAAA CAGCCAGTTO COCAGCACCO TTCTCOCCAGC TCTCCAAAAT CAGTAGCAAG TCCTCCAAAAA GITGICACIG AGTOGICCCC TGCTGGITGG GAGTGAAGAG AATCCAGGCT GGCAGAGCTG GOCGCAGACA GITTOGGACGA AACITTCAGAG CCCAGGCAGT CCCTGAATGA CCAGGCCAGT AGAGGAGGAG GAGTCAAGGG AGCAGGGCAG CTCTACCAGG CAAGGTGTTT CCCCAGCATA

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5 S GAGATGATTC TYTCTTOGCC CTGGCCATCT CGGGAAGCTT GATGGCAATC CTGGAAGGGT GOGAATOCTG CTGCTTCAAC CCCAGAGCCT AAGAATGGCA GCCGTTTCTT AACATGTTGA CARCCIAGAA ICCIGICACA CIACAGICAI TICTITICCI CICICIGGCC CIIGGGICCI AAAAGGTATA TATGCATATA TCTATATATA ATATGACGCA GAAATAAATC T TTAATCTCCT TTTGTGAGTT TGGTGGGGAA GGGAAGGGTA TATAGATTGT ATTAAAAAA

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 224:

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LENGTH: 2517 base pairs

SEQUENCE CHARACTERISTICS:

5 S GACAGIGGIC AGGGAIGCCI GGAGGCATAT ATCCAGCIGC CACCAAGGGG CACIGITITGI TITIGAATOGA AGAGGICAGT TIGTICCIGG CICICCATIT CIGGCCICAG TIGTCIACAG TCTGGTTGGG COGTTCTCAC CICGCCIGGC ACTIVACCAC ACCCIGGITT INIVIAGCCG CCAGCICICI CCCAGCICIC TITICCCITCI GOSIGICITI GCIGGGAGGG GGCIGIGITG IGAGCCCICC CAGAGGAAGT TOTOCAGAGT TOACCITTOC CITITICCITG AGITGIGGIG AATGCCCCAC CCTYTIGARAG GCTCAGCCTC CCATTOTOCA GTGCTTGGGT TTGGAGCTTA

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INFORMATION FOR SEQ ID NO: 225: Ε SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2424 base pairs TYPE: nucleic acid
STRANDEDNESS: double TOPOLOGY: linear

(X SEQUENCE DESCRIPTION: SEQ ID NO: 225:

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CCTGTGCAGA AGGAGAGGAA GGGGCATTAA GAGATGAAGG

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ATTACTTATO ATCIGIGICI AGGAGAAGGT

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TECTECACIT COCATGCCIC TAIGITACCC

GCCTCCTCCA CTCTATAAGC AGTCATCTTG

GGAGACCGGG

CATTICIGAA TAAACATITG TIATICCINA AAAAAAAAAAA AAAAAACICG AGGGGG

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	TIGTANCTAA TCGAGGATTG ATTCTAATGA CAGAGTCTTT CAACACTTTG CACATGATGT	09	CCCCCATGCT GTTTGTATAA GTTTTGCTTA TTTGTTTTTG TGCTTCAGTT TGTCCAGTG
•	ATCACGAAGC TACAGCTTGC CATGTGACTG GAGATTTAGT AGAACTTCTG TCAATATTTC	120	TCTCTGCTTG AATGCCAAGA TAGATTTATA GCCTTAATTC TTGGTCAGGC AGAACTCCAG
י	ITTCGGTTTT GAAGTCTACA COCCCTTATC TTCAGAGAAA AGAIGTGAAA CAAGCATTAA		AIGAAAAAA CITGCAICIT CAGTATACIT CCTAAAGGGC AATCAGATAA TGGATATGTT
	TCCAGTGGCA GGAGCGAAIT GAATTTGCCC ATAAACTGTT AACTCTTCTT AAITCCTATA	240	THAIGHAIT AAGAGTICAC TTTAGIGGCF TTCAITTHAR AIGGCIGTCF GGGAAGAACA
9	STECTCCAGA ACTINGADAT GCCTGTATAG ATSTECTCAA GGAACTIGTA CTTTTGAGTC	300 10	GOSTIGCCTA GCCCIGIACA AIGIAAITTA AACTIACAGC AITITIACTG TGIAIGAIAI
	CCCATGATIT ITITICALACT CTOSTICCCT TICTACAACA CAACCATIST ACTITACCATC	360	GOTOTOCTOT GTOCCAGITT TGTACCTTAT AGAGGCAGAR TGCCTCCGAR CGCTGTGGTT
<u> </u>	ACADINATAT ACCANTOTOT CITOGACCIT AITTCCCTTG TCHAGANAT ATCANGCINA	420	CITAITAICA AAAITAAGIT IACITGIAIA CGGAACAAC ACAAGAAAIT TGAITCIGIA
2	TAGGAGGGA AAGCAATATT COSCCTCCC OCCCTGAACT CAATATGTCC CTCTTGCCCA	480	AAGAATCCTC TTTAGCTGTG GCCTGGCAGT ATATAAATGG TGCTTTAFTT AACAGAATAC
	CAATGOTGGA AACCAGTAAG GGCAAAGATG ACOTTTATGA TOGTATGCTG CTAGACTACT	540	CTGTGGGGGA ANTAAAGCAC ACTTGATGTA AAAATAATTG TTTTATTTTT ATTGACATGA
20	TCTTTTCTTA TCATCAGTTC ATCCATCTAT TATGCCGAGT TGCAATCAAC TGTGAAAAAT	600 20	CTGATTGATT GCTATTCTGT GCACTNAATT AAACTGATTG TGATGACTTA AAAAAAAAA
	TTACTGAAAC ATTAGTTAAG CTGAGTGTCC TAGTTGCCTA TGAAGGTTTG CCACTTCATC	099	aadaadaaa aadaadaaa aada
25	TIGCACTOTT CCCCAAACTT TGCACTGAGC TATGCCAGAC TCAGTCTGCT ATGTCAAAAA	720 25	
3	ACTIGUATUAA GCTTTTTGTGT GAAGAITCCTG TTTTCGCAGA ATAITATAAA TGTAITCTAA	780	(2) TABPOBARATTAN STAB GED TT NO. 236.
	TGGATGAAAG AACTITITIA AACAACAACA TIOTCIACAC GITCAIGACA CAITICCITC	840	12) AND COLOMBIA DAY A COLOMBO TO
30	TAAAGSTICA AAGICAAGIG TITIICIGAAG CAAACIGIGC CAATITIGAIC AACACICITA	30	(1) SEQUENCE CHARACTERISTICS: (A) TANGET (1) SECTION (1) TANGET (1
<i>:</i>	TTACAAACTT GATAAGCCAG TATCAGAACC TACAGTCTGA TTTCTCCAAC CGAGTTGAAA	096	(c) ire: Mucrac acid (c) STRANDENESS: double
25	TITICCAMAGE ANGIGETICT TIMMATGOGG ACCTGAGGGC ACTOGETITIG CTCCTGTCAG	1020	IV) IUPULNAI: LINGAL (Li) PENITANE REPORTANI, CEN TO MO. 226.
3	TACACACTCC CAACAGITA AACCCAGCTC TAATTCCAAC TCTGCAACAG CTTTTAAGCA	1080	ARRESTANTA REPUBLICAN OF PAGE 1901 TO THE TOTAL TANKS OF TAIL OF TAIL OF THE TAIL OF THE TAIL OF THE TAIL OF TAIL OF TAIL OF THE TAIL OF T
	AATGCAGGAC TTGTCTGCAA CAGAGAAACT CACTCCAAGA GCAAGAAGGC AAAGAAAGAA	1140	אואואסמאני פאואאורופן ווארטוורופ זורורופטא ופרארוכארט שפרפסואיה
40	AAACTAAAGA TGATGAAGGA GCAACTCCCA TTAAAAGGCG GCOTGTTAGC AGTGATGAGG	1200 40	TAGOTGACAA GAAAACAAAG ATCITATTCA AAAGAGGICT TACAGCAACC CAACGICTCA GAATAACATA GAAAAAGA CACTACAAGAAAAAAAAAA
	AGCACACTOT AGACAGCTOC ATCAGTGACA TGAAAACAGA AACCAGGGAG GTCCTGACCC	1260	TETRICIPETE CANTENDENTY ABGRANCECCAC CACACGUSTY GINSTOTICAGA
45	CAAGGAGCAC TTCTGACAAT GAGACCAGAG ACTCCTCAAT TATTGATCCA GGAACTGAGC	1320	TOLINGUES CHESTER CHARLES STATE OF THE STATE
}	AAGATCTTCC TTCCCCTGAA AATAGTTCTG TTAAAGAATA CCGAATGGAA GTTCCATCTT	1380	ACCACABATA TCACTCTTT ATCACTANG TGGAGAACAA CAAGAATTGT GACAACAATT
	COTITICAGA AGACATOTCA AATATCAGOT CACAGCATGC AGAAGAACAG TCCAACAATG	1440	GOTTOCCACT GOAGINGAAC AAGGAAGGAA CTOGGINGGOTT TIGGAAAATGO TIGGIATATICO
20	GTAGATATGA CGATTGTAAA GAATTTAAAG ACCTCCACTG TTCCAAGGAT TCTACCCTAG	1500 50	ANCACHTOTH GAARMERICAG MUTCACHARCE ACTUMICACHAR INCOMARCA DAIRCTEATHS
	CCGAGGAAGA AICTGAGITC CCITCTACIT CIAICTCTGC AGTICTGTCT GACTTAGCTG	1560	CTGCCCGAR ANTIGCAGTT CCTGAGCTGG ATGGAAAGAC ACCAAAGATG TACAGGGGTG
۶,	ACTICAGAAG CIGTGAIGGC CAAGCTITIGC CCTCCCAGGA CCCTCAGGTT GCTTTAICTC	1620 \$\$	CEBBBBBBC CHEBACTER CAMPINGBBC CHINESTEED CAGEBBRETG CCCBBBRTTG
3	TCAGTIGIGG CCATTCCAGA GGACTCTTTA GTCATATGCA GCAACATGAC ATTTTAGATA	1680	GACTACOTOR INTERTIESCY CROCATOSCOT GEORGISSAA ANCCOTOAN
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2 INFORMATION FOR SEQ ID NO: 227:

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CICTICTION TGAGGGCTAN GCCGGGTGGC ANGITTICCAG GGAANCIGGA AGCGTTIAGA

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ATCCACCCCT TCCTGGTACT TCCCTGTCTC TACAGCCCAG GCCTTAAAAA TGGTATTTTC TICTOTOCIO TIATIGAGAS CACASCOCAS CIGOSCOTIC CATITAGOIS GOICAGOSTO ATATATETTE ACTORGITEA TOROGACETE CETTOGTOCE ATECATACAT CEAGGITGAA

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(A) LENGTH: 1336 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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EX. SEQUENCE DESCRIPTION: SEQ ID NO: 227

GSCTGCCASC TCTCCTGGGT TTGAAGGATG CGGTACASCT GCTTCAGCTG AGCAACGATG TIGOATICAC AATTACTOGG AGGCAGOCAG GGGCAGTIYGC ATOCTGGGGG TOGCTGCATG 120 6

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TTATCCTTGA TOTCTOGGGT TGAGATCTOC AGGCGGACAC TGCCACTATC AAAGGATCGT

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SEQUENCE DESCRIPTION: SEQ ID NO:

228:

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 2043 base pairs 2

(2) INFORMATION FOR SEQ ID NO:

228:

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SEQUENCE CHARACTERISTICS:

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AAGATGAGTT TECTTTECAA TOGTTTECCA TETOGCCATT CTTCCCCAAA GCATAAGTAG GTGAAATCAC CAGAAAACAT CTCGTAGATC ATCCGAGCCA CTACTGGAAT GACCTGAACC

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ATCTCAAACG GTGGCTGCTT CTCTATCTGT CCTTTCTGGT GGGCAATGAG ATCGCTAAGG

AATGITICCA GACAAAATAG CITGACCITC TITTGICICT CAATCAGGIT GGGAGCAACA 420 360

AGTGATGGGG CACATGGCCC AGACCAGTAC ACCTTGCACT GGCACAGYCT GATGGCATAA

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ATROCATORC COCTORCCTC CAGGATCAGT CCTCTOTCCA TORCOTCCAG CAGCTTGCTA

GTGAACAGCT TCTGCTTCTC ATTGGTAATA TGCTCAGGAC CTGGGAATTT GACCTGCTCC AGNOTIGAÇOG GACCAMAGAG CITCOTOCIOG ICAGGOATOG GACCCAGGIC COCATAGAAG

AGTEGGEAGE CETGAGGGTT GETCACGGTC ATGGTCCTGC CCGTACTCCT TCCCACGGTA

CTGARACTTC ATGTCCAGGT CAGTCATTOG GAGAGAGCTG ATCCACAGTT CTGGAGAGCT ATAGAAGGRC TGTATAGGTG CCTGGGGHAC TICCATCICC AGGGGTICAG TITITGGGCCA

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CAGICIGACC CIGCCCCTA CIAAAGCCIC INCICICAGC ACTITICCCCC AAGICCIIGI

GCTIGGACTIGG TICCIGGAGCTIG GCTTTCCTTTAC CAGAAAAAGCC TICAGCCTTTCC TICTIGGAAGCA CTCTGCCCTC AGGGTCAGCT GCCCAGACTG GGGGGATGC AGAGAGGCAG GTGGGCTGTG CADODCTOOG TOOTGAGCAG CTGGTTTTTCC TOCAGGAAGG TTGGAGCAAG CAAAGTOOTT CCAGYTCAGG TCTGAGACCC GTGYTGAGTA AAGGTCTGAG CAMCGACCGT GCCCTCTGCC CAGAGGCCCA GGGTCCAGCA GCCCGGSGGG AACGGGTGCT GCCTSTTCCT CCAGTTAGCT CICAMETECE CITECETBAG CICCITECEA AGGACICETO GICACIGEET GETOTGEAKT CCCGCTOCCT TOCAGCCTOC ACTITICAACA TGCTCACCCC CAGCACAGTC CCACTGGCCC GALCTCAGAC CCCACCCACA CTCCAGATCO AGACCCCTOC CTCCCCCCGO CAAATOTCCT TEAGETOGIC CETTECTION GICCIGGGG ACCIGCIGGC GGCCTCTICC IGGGAGCCAN

TOCOCOUTTO TODOCAADOO GGAADOOCTO CTITAADOG TOTOCTITICO CAGTODOGAG

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ATCCAGCTCA TCTTCCTCAT CTTCTTCATC CACATCATTA TCCTTCTCAT CCCAGGGAGC

960 900 840 780 720 660

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AGACCCTOTO GATCCTOSOT TAATGATCGA SCCCTOGGGC TGAGGGATOT CACACACTTO

CACTRECTICE, GGSCTGCAGT TOCCCACACT GCAATTGCCC ACACTRGCTG GCGCCATGGG

ATGITICAGGA AGGGGAAGGI GICCIGGAIG GGAACAIGGI GCIGCGACIG

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ICTIGGTCTT GGAACICCCT GGCATIGGGA ACAGAGCATT TCCAGCATTT GTTGTTGTTG

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960 900 840 780 720 660 600 540 480 420 360 300

CCAGGCTGGG AGGGTTCCTT CCCTAGCTCC CCATCTGCCC CCGCTGGTGA GAGTTGGGCT NGCCEAGETG GGGGGECEGG ATGGGGGETT CTETETETG GAGGGGTGCA GGTGCCETEC CCTOGTGAGG GGAGGTGCTG CTTTTCTGCC CCACCTGCCG GCTGGTTCCA GCAGCGCTGG TITITICATIT TITICITITCCG TCTTTCTTCT TGAGTTCACG GITCAATATI GCCTCCTCGC TITAMATTIC ACTUATITIG TATAMACCCA GCAGGCIGGI GITIACTIAG CCCIGIAGCI AACTIGETIG AAGGIGGGIT CIGGEIGEEA GEEAGIEEET GGACAAACIE TECTIGEEEET

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	177		•	-	
	TITIACTCAC CTAACCCTTA GAAAATGAAT GITAGAAGOT GCCTGCCCGAG GCGGGACAGA	1140	ASSESSED OF THE PROPERTY OF TH	ADDRESS CHICAGA CONTRACTA	084
	GTOTTHECT GOGCHOGAGA AGGENCTICCT CAGCOCTICAG AGTOCOTTICC TGCCCCACCG	1200	TOTIOGOGIC CONCOUNTS CUITORING CIOCURANO SITUADOS STATEMENTS	COCONT APACACACAA ACCTICABAAT	0 9
5	ATACTGGCAC TTTAMANAGG ANGCTGACCG CACAGTGTCC AGACGAATTG GCCCCCAGAA	1260	3		į
	GATGGGGAGT TETGTECTEC CETTETGTS CTGCGTGACC TEACCEAGCE TAGGAGGGAG	1320		:	
9	GTOCATTCAG GGTAGATTTG CCTCTCATTC AAAGTTCTGG GGCTTTGGGY GGAAAACAGC	1380	(2) INFORMATION FOR SEQ ID NO: 230:	-	
2	CAGCITITGGC GCITOTIGGGG AGACITCCICC AGACCAGAGA CCCCAGAAGG AGACAGAGCC	1440	(i) SEQUENC		
	TOCCACATOC TOCCACOCCA GOCCOTIOGOC CAGGOTGATT GAACTGAGAA TITIGOCCACA	1500		711	
15	ACCANATICA TOCTGOCTGO AACCAGAGGC CAGAAAGCCT GOCCTTGTCC CCATGTGGGA	1560 15	(C) SIYANDELWESS: GOUDIG (D) TOPOLOGY: linear	- -	
	GOCCITOTOCT CAGCOCTOTT GTCCCCTTGA GCTCAGTGAA TTCCCACCAG GTGCCCACAG	1620	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 230:	ID NO: 230:	
	CTCCTGGACT TCAAATTCTA TATATTGAGA GAGTTGGAGA GTATATCAGA GATATTTTG	1680	AATTGTGAAA TATTAGAATA TTGTTACTAT TTGACCCAAC TCAAAATCTC CATGGGAAAA	CCAAC TCAAAATCTC CATGGGAAAA	09
3	GAAAGGAGTT GOTCTATGCA ATGTCAGTTT GGAATCTTCT TGAAAGTTTA ATGTTTTTAT	1740	TACCTGTCGA TACCCACAGT ATTGTTCAAA ATAATCAGAT GCAGTATCAC AGCTGTGTCA	ICAGAT GCAGTATCAC AGCTIGTICA	120
	TAGGAGATIT AAAGAAAATA AAGGICTACA ATATCTITAG GITTITITIT TITICCIGITT	1800	GACTETAGTA CCAGTTGGGC ANTCAAGGCA CAGCTAAAAA	TAAAAA TIGAAAACAA AGAICTOGAC	180
25	ACCICIALAA CTGACCACAT GOCUTGTCTA TCAGGATGGA GOOTGTCCAT GTTCTCCTCT	1860 25	AACAAAACAG CCAAAGGTGG GOGTCAAGAA GCTCTGACGT GTACCTAGCT GTAGAATGCT	GACGT GTACCTAGCT GTAGAATGCT	240
	STCTTTAGGG AGGIGARAG GAGATGGSGG RAGGGGTGTT TITTTCTTTG ACTCCCCTCC	1920	атесасаст ессастета стетесатат сеасовалала стесасаса сеселетт	HANANA CTGCAGAGAG CCCCAGTCTT	300
90	TITICIDACAG AATGITGCCA CCACTGCTTG AGTGGGCTGT GTTTGTTCCT CTGTCCCAGC	OE . 0861	CANCICTOGT TEACCATEAG CTCTGTGTAA GCAGGAAGTG AAGGCTAAGG CAGATTTAAG	MAGTG AAGCCTAAGG CAGATTTAAG	360
3	TICTOTICIA GAAAATAACA TICTIAGGG AACTCAGGCT AGTGTCAGGG TCTTGGTTTG	2040	CTCTGAAAGC ATTCCACAAC ATACACAA ATCGTGCAAA GCATTAAGGA AATCTTGTTA	GCAAA GCATTAAGGA AATCTTGTTA	420
	555	2043	CTGCTAAGTG TTGCTGACCC AGGAACAA		448
35		35			
•	(2) INFORMATION FOR SEQ ID NO: 229:		(2) INFORMATION FOR SEQ ID NO: 231:		
04	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 540 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 407 base pairs (B) TYPE: nucleic acid (C) STRANDEZNESS: double (D) TOPOLOGY: linear	82.1	·
₹	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 229:	U	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 231:	ID NO: 231:	
	TAMAMAGAMG COGGAGAMTC TOOSCOTCOC TCTAGAGATC GATOOOCTAG AGGAGAMGCT	09	GTATGCTGCC CCAAACCAAT ATGTGTGGCT GCCTTTWACC TGACTTCTCC AACATGTAGC	TWACC TGACTTCTCC AACATGTAGC	09
20	STCCCASTST CGGAGAGACC TGGAGGCCST GAACTCCAGA CTCCACAGCC GGGAGCTGAG	120 50	CCCAAGAGGA GGCCTCTAGA CTRAGGGAGG GGCTGGTGAC CCAGGTGTGG	ISTIGAC CCAGGTGTGG TGGGGCTGCA	120
	CCCAGAGGCC AGGAGGTCCC TGGAGAAGGA GAAAAACAGC CTAATGAACA AAGCCTCCAA		TGARACTACC AGAGAGACAG ACATTCTGGA ACTCACCCTG GGGGATCCAG TGGATCTGCC	CCCTIG GGGGATCCAG TGGATCTGCC	180
55	CTACGAGAAG GAACTGAAGT TTCTTCGGCA AGAGAACCGG AAGAACATGC TGCTCTCTGT	240 55	TATIGOTOTICG TOCACOCCAG ACCTOTGAGA TOTTCCTCAT GAGGATGCAC TTOTGCTTCT	CTCAT GAGGATGCAC TTGTGCTTCT	. 540
3	GOCCARCITY ATCCTCCTGA CGCTCGTCTA TGCCTACTGG ACCARGTGAG CCTGGCACTT	300	GCAAOTATTG CTGCAGCTTC ATAGTGACTC CCACCAGCAC CAGCAATACA GYTAGCTACC	AGCAC CAGCAATACA GYTAGCTACC	300
	CCCCACAACC AGCACAGGCT TCCACTTGGC CCCTTGGTCA GGATCAAGCA GGCACTTCAA	360	TOTGGCCTTG GATCTCAGCC AGCATGGCTG GGAGAGGGAG CARCTGGGCA TGTACCCTAA	GGGAG CARCTGGGCA TGTACCCTAA	360
9	GCCTCHATAG GACCAAGGTG CTGGGGTGTT CCCCTCCCAA CCTAGTGTTC AAGCATGGCT	420 60	ATSCTIGITAC CAGGAAAGA CTCCCAGAGT GAAGACAAGT AGGGACT	CAAGT AGGGACT	407

(D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 830 base pairs

STRANDEDNESS: double

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(2) INFORMATION FOR SEQ ID NO: 232:

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SEQUENCE CHARACTERISTICS:

WO 98/39448

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ઝ 25 20 45 8 ઝ 5 8 55 8 AACAAATAAA GCAAAATATA ACATGCATTT CACATTTTGT CTTTCCCTGT TACGATTTTA TAGCINCOAG INCTITUGAA CTATAAAGAT GICACTACIT AACACACATA CCTTATOITI GTATTYGATT TCAGGCTGCT ANATGGGCTC ATTTAGCATT CATTCCTTGA TGTAGACATT ATAAATACAT CAGCACTGCA GCACACGITT AAGGITIGCCA CGGACAAGGA TCACACAATA ATAGCAGAAC TOTATGACAA GTTTAGGTGA TCCTAGCATA TOTTAAATTC AAATTAATGT TAGAATOOTO TITTITTATIG TOTTOTAAGG ATATGGATGT TOCCATAACA GCAACAAAAC GITTATTATT TITCIGITAA AAAATIGIGA AAAGITIGIT TIAGCIAGAT GATATIIVIAA GAGAÀCACTO TAGTTCOGTC TOCTCACAAG ACCCAGAACA TIGATCAGTT TITIGTTGTTG CTACOGRAAC AAAAGAGOTG AAAGAGACCC TTTTTTTATA CTTAATOTAC ATATATTGAC ACCAGANATG AACTTGGCCC TAGACCTAGG GGATAAGCAA TGTTCTTTAT GTAGCCAATG CCAGAAGAAA GACCAATCTA GAATATOGAA CTCTAATCAC TTCTAGTATT TCAACTTCCT 2 AATAACTYOG CYTCGGYTAT CCATCAAATG CACACTTATA CYGTTATCTG AAAACAGATT AACAACAACA AAGAAACTGT CTATTTGAGT GAAGTCATGC TTTCTATTAT AGCAACAAA ACATITCATA AATATCACTT GATAGACTGT AAGCACCTGC TTAACTTTGT IOTITIGITT TOTITIACAC TCAGTATAAA TCAGGAGAAG TTAGCCAACC ATCTAGCATT AAAAAAAAA CTGAATAGCA TTCTTTCCAG GNTAACTAAT AAAGCAGACA TGCTAAGCCT INFORMATION FOR SEQ ID NO: 233: Ê (xi), SEQUENCE DESCRIPTION: SEQ ID NO: 233: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear (A) LENGTH: 932 base pairs 600 830 780 720 660 540 480 420 960 900 240 180 120 180 120 8 8

> 25 20 5 ᅜ TACATTITICT CATTAGGGTT RIGATGCTICA GTATCTTTCC AAGTGCCAGG CAGRGCTTNC AATTIGACAA AGGGIGICAT AIGCITTICCT AACCIGAWIT GIATIAACAT TCACAGAGGC GGAAAATTGA TACTITIAAA GCATATICTT CIATGAGCAC AGGICCTCCT AGTGAAACTI TCACTATCAA GAGCCIGCAG AGCCATTTIC CAGACCIGIG ATTGCCCAGA ACACATAGIC GCATTGTGGA TTAGCCTGAG GCTTAAAATC AGATGCATGT CTGGTAAGAT GACCACTGTC AGGITICCIA AACIATAAAA GCAGATIITIG CITITIGITIG TIAAICATAG GCATGGCCGA TCCCAATTTA AAAATGCAGA CITITICIDAT CANACATACC ATTITITIOTA TITICACAACT ATAGACAGIC ACTICIOCAG GCTGTGTATT GTGACTATCA GCATTCTGGT GCAAATGAAC TECTGARTIG GARGAGGAAG RACTICIGIT TACAGAAAAC YGTATIGITA TATATGICAG GCTTTCTAAA ATGRANTCAG TTTCTAAAGT GAAACATGCA ATATTTATGC TITITIGAGCA AGAATGOCCAG AAATAGOCTT CATTICTACC CIGCAAAATA AICCAGATCT CCCACGITIC TAATITIGGAG CAAATCTAAA AG ACTOCTITAT CCAAGAATGC TGAAAAATAC TGTTCTATCC TITTICTCCAT CATCGACTGT TCTGACTGAC 900 660 60 540 480 420 360 300 932 840 780 720

2 INFORMATION FOR SEQ ID NO: 234:

30

E SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (A) LENGTH: 2786 base pairs STRANDEDNESS: double

SEQUENCE DESCRIPTION: SEQ ID NO: 234:

(D) TOPOLOGY: linear

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S **£** 6 S TTAGCAGGGT GAGCTGTTAA AACAGCACAC ATCTCTCATC CCCTCTTCCT TTATTCCCCC TEAGRECECE TECENACEET CEATHIGGET CICAARGGIG CICACTIGGT TGGAAGCAGG CTGGGTTTCA GAAAGGAAGG ATATATGGGG ACCACCTCCC CCTTCTTTGA TCCCAGCATC CAAAATCACT ACAATAGCCT AGTGCTTTTT TOGAAGCCTT TTTAGGGAAG AATGTTAGGT CATGGGATCC CTTCCATAAC AGGTACTTTG AAGGCAAGAC ATAGGGTTGA AGAAGCACAA IGITITIGIG TIGIGGAACC IGAGATICCT TATTIATIAA CAGGAAGICI GATTITITIT TCAGTOAGAC GATGAGAAAA GTCCCAGGCT AATGGCAGAA ATTTGCACTT TGAACATGTG CICCOAAIAG GGAGGGGSCI GCCCICIACA GICICITIGA CIGIAAGACA GGGCICIGIA CCAGCCTCTG AAATCATAGC TCTCCAGTGG CTTTTAAAGA AAGCTGGTCC TCAGCACTAA TTTTCGAGTC TTTCTTCCTA TATTTTCTCG GCCTCGGAGA GAGAGATTAG ATTATTTTCA 300 240 180 120 540 480 420 360 600 8

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	461			462 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	**	
	TCATOGINAC TNGTATGCTC TTTGAGATTT TTACAGTGTT GAAACTTAAG AATTTTGAGA	099	4	ACTACCATOC ACATAGACA CITIGAACCTO CITIGIACCT GITIGGGGAA AAAGTATAAT	2460	
	COCICACCAC COTTOTICAG ARCTAART ACACATAGAT CATTOTITOT TOTCAATITG	720	Ü	GACTETACTA CCAATCTAAC TAAGATTATT ATACTCTGGT TGTTTGAAAT ACCATTTTTT	2520	
2	THICHTING THITHTING TOCHAGGA THOCHIAGAT THOCHINGG GOGGANCHCT	780	2	TCTCCTTTTG TGTTTTTCCC ACTTTCCAAT GTACTCAAGA AAATTGAACA AATGTAATGG	2580	
	GACTOCITICS TITTIFOTITIC TITACHATIC ATTICAGETC CENSITIAGIS	840	•	AICAAITITAA AATAITITIAT ITICITIAAAAG CCTITITITIGC CTGTTGTAAT GTGCAGGACC	2640	
	AAGGACACTO CTOTTAGTGA AGGAACAAAG TCTATGAGTC CTAAAATTTT AAGTCAAAGA	, 006		CITICICCITY CATGGGAGAG ACAGGTAGIT ACCTGAATAT AGGTTGAAAA GGTTATGTAA	2700	-
2	AACTGCTCT GTTTCCCCTT TAGTAACACT TCTGAAGAGG AAAACTTCA ATAGCCAAAG	096	2	AAGAAATTA TAATAAAAGG GATACTTIGC TTTICAAATC TTTGTTTTCT CTTATTCTAG	2760	
	TTARIANICC TATATANIA TISCITIOSC TITCACCIAA ANTICIOSSC AICACANITI	1020	Ö	GTAAGGCATA TTAAAAATAA ATATGT	2786	
5	CCTITOGGATA GAGGITGIOT TGGGGAATAG ATTGCTTATT CCTGTTCACT GGAGAGAAAA	1080	15	•		
	COTACTOTIT TIGTACAAGG TCATACCOCC AGAACCCCCA AATCCTATIT TGGCTCATCT	1140				
9	TCAGGIAAAG AGTAATTCCT ATCCTGTGTG CCTCAGAAGC TAGAATCGAA GOCTTACCCT	1200				
೧	ATTICATIOTY TATTOTCACA AATOCATGAT OSCICTIOGA AAGAATGAGG TITTIGCTIGGA	1260	3			
	aaaaaaaa agaacagtit gtotticaca aacatgoctt atcaattitt tcaaagaatt	1320		(b) TIFE: INCLEAL GLID (C) STRANDEDRESS: double (D) ADDITION: Illustration		
22	CITITITICC AMANGNOSA GINACAMANT GICATITICIG MANGNOSCIT ACTITININC	1380	25			
	NACTRATOTIC AGCUTTITICOS ATOCICAGOSA ACAGAGATO AGACACITAC ANTICACIACT	1440	,	(A.) SEQUENCE DESCRIPTION OF SECURITIONS OF SECURITIONS AND SECURITIONS OF SECURI	. 9	
9	CICAAAIGCG CIAITIGITHC ITITICAGAGT GITIGCAGAIT IGCCAITITCT CCAIAAIAIG	1500	, , ,	COLICIONE DILICOGRAD REPORTED INTERPRETATION OF THE PROPERTY O	120	
≥	GOGATAGAAA ATOGAATAAA GATAGAAGGG ATOTAGAATA TGCTTTCCTG CCAACATGST	1560		TUTIOTICA TOTTUNAMEN TIACCITAGI GAMENIGIGI TICACCITA		
	ITGGAGTOGA CTTTGGTATA TTGACTAGAT TTGAAAATAC AAGATTGAYT AGATGAATCT	1620	۳	TACCTGCATA ATCACACTA TCCATCTATT CAAGTGATG ATCTGTGGGA TACTTTTAAT	180	
35	ACAAAAAGH TOTCCTCCTC TCAGGTCCCT TTTACACTTT TTGACTAACT AGCATCTATA	1680	35	GREGOTCHCHA ACCHARACT NGATOTHANG AGARTGATGA ANACTIGGAC CCTGCAGAAA	240	
	THECACACITY ACCITATING NEACACITIAN CETTIONERS OGNABATITY ATTITICAGIG	1740		GGRIFFUCTI TRICIARLIST TUMPANAMA GENEROUMA. IIIIIninus munomanama.	360	
	STINGTONIC AGNINITITA SCCACCINCA CANANGCANA CIGCATITITI ANAMAICITIT	1800	, ,	INTITUTE ATAIONIC TOTAL COLUMNIA CONTINUE CONTIN	420	
2	CTGAGATGGG AGAAAATSTA ITCTCCTTTC CTATACGGCT CTCCCAACAA AAAAACAACT	1860		THICKNOCK ANCIETTION CALLETONS INDICATE LINGUIST TOWNS OF THE CONTROL OF THE CONT	, a	
	AGTINATICE ACTANTINGA AACTIGCTOT ACTITITICIT TICITITHOS GOTCHAGGAC	1920	•	Addictive GAMECHANE ANDALANCE CITECIAN	}	
5	CCTCTITIATA GCTACCATIT GCCTACAATA AATTATTGCA GCAGTITGCA ATACTAAAAT	1980	45			
	ATTITITIMIA GACTITATAT TITITOCTITI GATAAAGGGA TGCTGCARAG TAGAGTTGGT	2040	Ū	(2) INFORMATION FOR SEQ ID NO: 236:		
S	GIAATTAAAC TATCICAGCC GITTCCCTGC TTTCCCTTCT GCTCCATATG CCTCATTGTC	2100	ç	(1) SEQUENCE CHARACTERISTICS:		
3	CTICCAGGGA GCICTITIAA TCITAAAGIT CTACATITICA 1GCICTIAGI. CAAAITICIGI	2160	3			
	TACCTITITIA ATAACTCITIC CCACTGCATA TITICCATCIT GAATIGGIGG TICTAAATIC	2220		(D) TOPOLOGY: Ilnear		
22	TGAAACTGTA GTTGAGATAC AGCTATTTAA TATTTCTGGG AGATGTGCAT CCCTCTTCTT	2280	55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 236:		
	KONGGITIGCC CAAGOTTOIT TTGCCTIAACT GAGACTCCTT GATATCCTTC AGAGAATTTA	2340	4	AGGATGAAGA GGAAATTAITC TCTTGGATTG CTCTCCAGGA AATCCTTCTC TATACTTTAA	09	
ç	GOCAAACACT GOCCATGGCC GTGGGAGTAC TGGGAGTAAA ATAAAAATAT GGAGGTATAG	2400	ç	ANGCICTIOT TETTTTETAG GARTECAATG TGCTGATTOC TGCTAACAGT CAGGGTACAA	120	

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20 5 25 5 35 30 45 6 S S ş TETTITIANG GINCAGCAGG GANGANCIGG ANACICAGAG ANAGANACIG CECTICENIC GTAACAGTGA ATAATTTGTA AAGTTCGTAT TICCCCAACCT CITTGGGAAT T TGAATAAAA TAAATAACHA AACAATAAAA GTTTATTGAG CCACAGTTGA GCTTGGAAAG AUGGACATIC CICCICITICG TOCHCITATC TAATITITICT GATAGGGAAA CAAATICITIT CARAGICATE AUGGSTITIG GARTISTITE GARTATITE TENTETIC TEXTECCTES AAATACATOT GCAGOTGACA ATGAGAGARG AAACAGAAAA TGTOATGTGA TGTOTCTOCCO TTAAGGTGCT AGAATTGGTA TGAAGGGTTA ACTCAAGTCA AATTGTACTT GATCCTGCTG CCTTCCACCT CCTCCATTTC TICCACCTCC TCCGACTGTC AGCACTGCTC CACCTCTGAT 3 TTTTTOTCAA ATGCNGCAAG AGATAACICT TITTANGAAG TAGCATATGT GAACTATAAT TITATGAGCC CAGGCGACGG GCAAATGAGA ACAGCAACAT ACAGGTCCTT TCTGAAAGAT CTGCTACTGA GAGATTACCT GGGGCAATTG ATGITATCGG TCAGACTATA ACTATCAGCC GAGTAGAAGG TACAAAACCT GACTITACTT CICCICCTIC ITIOTICAAG ACTOGGCTIC CACCGAGCAG TOCCATGAAG AACGATACAG ATACAGGGAA TATOCAGAAA GAGGTTATGA GCGTCACAGA AGTAGACAAC AATTITAGCA AACCACCTCC GTTTTTCCCT CCAGGAGCTC CTCCCACTCA GAGCCTGCCC CTGAACAGGA GAGCACCGAA GCTACACCTG CAGAATAGGC ATGGTTTTGG CACAGGAGAC ACAAACACAA AAAATCTAAA AGAAGCAAAG AAGGAAAAGA AGCGGGCAGT AGACATAAGT CTTCTCGAAG TAATAGTAGA CGTCGCCATG AAAGTGAAGA AGGAGATAGT GCAAGTCGAG AAAANGAAGA ACGACATAGA GAAAGACGAC ACAGGGAGAA AGAGGAAAACC AGAAAGTGGA CATTCCTCTG GTTATGATAG TSGTTCTGCA CGTGCATTTC CATATGGCAA TECHECACE GUITTICCIE CIECHECHGG CUCICCHEET CENTEITHA TACCHACHAT TOTTUAGAA ATTIACCITA AATCITGITC TOTTTGITAG TAIGAAAAGT TAACTITITT INFORMATION FOR SEQ ID NO: 237: E χ. SEQUENCE CHARACTERISTICS: TITIGGGACAT TOGGANIACC CAGCCAACTC TCCACCATCA ATGTAACTCC ATATTAGTAC CAGAAGTAGA TACTATAAAT CITGITATIT TICIGGATAA SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 1286 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 480 420 360 300 240 180 120 540 720 660 600 8 900 840 780

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INFORMATION FOR SEQ ID NO: 238:

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GGTTTTICCNC ACCITTTIGG TIGGGC

CCATTACTOC TOTOGACACT CTTOGCTTRG TATWITTAGG GOOGNTCCTT ACCITITITE

1260 1200 1140 1080 1020

ATCACAAARG GCAACCAARG GOCCCCTCTT ARGGCTTTGA GGATTAAAAC TAGTCTTTAT AGGGTICCAC TIGGGCCACA GITTITITIGI TAAICAAACA CCACICICIT AAGROGCIGC

S

CAGOGNATICT ANAGAGETIST STEAGETISTS TACATACACA GATTATICTICA GAANAGGTICA

aaaaataaaa agacagcaat gactttatat ccaagaaagg aatgtgaatg agtcacttaa

SEQUENCE CHARACTERISTICS:

20

Ê SEQUENCE DESCRIPTION: SEQ ID NO: 238: (A) LENGTH: 734 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

30 25 45 6 ઝ 8 GIGITECTICG GEOTEATICET GRACIGIGIG GIGAEGIECE CIATOTIGET GGIGGETETG CIGCIGCCGA AGCIGATICC CICCOGIGCA GGCCGGGAGT GGCTGGAGCG GCGCCGCGC ATTOCCACCOC AGAAGGACCA GCAGAAAGAT GCCGAGGCGG AAGGGCTGAG CGGCACGACC CAGATOGAAC CCGTGTGAGG TGTCTTCTGG GACCTGCCGG CCTCCCGGGC CAGCTGCCCC CCTCTTTGGC CGAAAGGTGA GCCCAGCGCA TCATATGCTC TGGCTGGAGG CATCTCCTTC GEAGAGETOT GECAGEGEET COTAEGEAAC GTOGAGTAET ACCAGAGEAA CTATGTGTTE ACCATICCOOC CCTOGRACIAC CTTCGTGGAC CAGCAGCGCT TCTCACGGCC CCGCAACCTG GIGGTEATEG GETECEAGGE TGECTTECAE CAGATTGAGG CTGTGGAEGG GGAGGAGCTG ecementer acerdacias recosseres occonenter assistensas asceaeceps OCTOTOTTTT TOGGEOGETG TTAACATTOT CTATOTOGGO ACCITOGAGT CCAAGOTTOT CAAGCCCOOG GAGGGATICCC GCCTTTGAAA ATAAAGCTIGT TATIGGGTGTC ATTICAAAAAA ACCCCTOCCC ATOCCTOTICC TOCACOGTCT GCTGCTCGGG CCCACAGCGC CGTCCCATCA алалалалал алла 360 240 180 120 540 80 420 9 734 720 660 60 60

2 INFORMATION FOR SEQ ID NO: 239:

SS

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 809 base pairs (B) TYPE: nucleic acid

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TCCARARTAA AAGAGTGAAT TTTTCATGTT AAGTTAAAAA TCTTTGTCTT GTACTATTTC

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1740 1320 1380 1440 1500 1560 1620 1680 960 1020 1080 1140 1200 1260 GOCCOTOTIC CCACOTOCCA GACACTACAT GOGTAGCTCA GOGGAGGAGG TOGGGGTCCA AAAATTITGTA GCCAGACCCC AGGIGCCTGC ICTCGTCTTT CTCTGGGTGG CCTCTGATC TENGCISTAC GROSTSATCC CTACCCGACG CTCCACCGCC GAGGCCTTCC AGATCGTCCT GETCHGGGG TIMETIGGG CACTGGGGG CGCACTTICC 1GGGCACGG CATCTTCATT GAGGCCGACC GCCGGCGGC ACAGCTGCAC GTGCAGGGCC TGCTGCACGA AGCAGGGTCC ACAGACGACC GGATTOTOGT GCCCAAGGG GGCCGCTCCA CCCGCGTGCC CGTGGCCAAT GIGCICATOT GAGARGOTIC COCTOACOTA COTOCACATO TGCCACAGOT GGCCOTOGGG CCACCCCACG AAGGGCCTGG GCCTAACCCC TTGGCCTGGC CCAGCTTCCA GAGGGACCC GANGGAGAT COCTOTOCAC AGGGCAGOC COAGGGCTO GGTGCTAITT GTAAGGAAN ARGOGOTOCA GOGCATCAMO GOCCTOTOTO COGGCOGITO GACTOTOATA GTOGGGGTOC AGIATETEAT GIGGGGGGC ATTGCCTTCT GGTCCCTGGT GACACTGGGG TCATCCTTCA TCCCGGGARA GCAITTCTIGG CTGCTCCTCC TGACCCGGGG CCTGGTGGGG GTCGGGGAGG TOGGODAGNOC CCACCOTOCC TTCCCOGAGA CTCCTGCTCT TCCTCTGACA GTCTCATCIT cosectedes encretance ecososetan recectoste totocenero ocetectosos CTCTGCACCC TTCCTCTTCC TGTCCCTTGC CTGCGCCCGT GGTAGCATCG TGGCCACTTA TAITITHATC TICATIOGAG AGACCCICCT GICCAIGAAC IGGGCCATOG IGGCCGACAI STOCCACCTG CTGGGTAGCCC CTACCTCATT GGCCTGATCT CTGACCGCT acecceana nececce cenentore cantroces cerenean reneorichi Greenveces ACATICAACA OTTICTICAAC ATCGGGGACA GTAGCTICTGG GCTCATCCAG ACCGTGTTCA TCTCCAGITA CATGOTOTIG GCACCIGIGI TIGOCTACCI GGGIGACAGG TACAATCOGA GACCAGCGGA TGACACCCGG CAAGGGGAGC CTCCCTGTGG AGATCAGCCG GCTACATTGC TOTCACOGO TOCOTOSOCTO TOTOGOCTOC GOCATTOCTO CTGCOTTOCO GOGTGOTÓT COGGATICCTC AGCATICTICT ACTITICACAT TCCGGTGGGC AGTGGTCTGG AGGENCEAAA GIGAAGGATA TGGETGGAGA CTGGEACTGG GCNCTGAGGG COTOGRAGOSC CACTCAGATT TOCCACCCCT GAACCCCACC TCSTGGTGGG GCTICTICGCA AGAAATCCTA GTTTCGTCCT GTCTTCCCTG GGCTTCACTG TETAGGAGIG GIGGCGTTC TGCTGCTGTT CCTGGTAGTG CGGGAGCCGC CCAGTTATTC CACCATCGCG CCCACTCTCA TTGCCGACCT CTTTGTGGCC TOTOCTACAT CANTETECTS AACTACATGG ACCGCTTCAC CGTCGCTGGC TOCACCCCGT CITICACCCCA GOCCTCCTGA AGACTIGTGGG T TOSACTICATE ACCTGCCTGA CCGGAGTCCT GGGTGTGGGC

2160 2201

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2040 2100

5 35 30 25 20 ᅜ 8 45 55 50 CATCHTTCCC CCTCCCATCA TACCTCCTCC TTCCTGGAGC CTCTGCCGGC TTGGCTGTAA NGACAGAAAA GCAGAAGAIG AGACICIGITT CATICACTIT ICCIAGGCCC AICCIGIGGT ຣ CAGCCICCIG GIGIGIATCA TAGGATTIGI TCACATAGIG TTATGCAIGA ICTICGIAAG TOOQACAGAT GOOGAGAGGA AAAAGGCAGA GATNGCCAGG AGAGOOGTOC AGGACAAACC TOSTOSCACT TACCTOGATA TTTCAGTOGG AGGATGAAAG GCGAGACTCA CCCTACGCGG TAATICAICA AIGITCIAGI TAATGICIAC CICAGCACCI CCICITAGCC TAATITIAGG GITTAGCCAG CAGCTOCGGC CTCCCCGGGC CCTTGGCATC CAACTTCGCA GACAGGGTAC AGAGAGITTO GOTCAGOGA AAAGTOTNOG GAGAAAGTOG GOTOCAGOCC CTOCAGGCCG TETTATOTEG TGATAGEACA AGTGEEAGTE GGATTGETET GTATTAGAGA ATAGTGTTTT CTARGECTIT TOGITTOGGT ATTATOTITIC GITTIGITAT TIGITGGTT TIGITGGCTTG ATATRIBITOT ATGRAARTA TAGCACIGAG GGCCCTGCTG CCCTGCTGGA CCAAGCAAAA GTTANGANGC COTOGTGGTG CACCATGACA TCCANCCCGT ATATATANAG ATANATATAT CTOTICIAGI TECGAAGEAG TITICACIOGA AGITOTOGAG ICCIOGITOC AGCITICOGA TRETERETT CICTECTECE CACCICICAC CETTGECETE TECATETECE TETECEGECE AGGITGCCCA ATTITGTTIC TICAATTITA CIGGITACTI TITITGTACAA ATCAAICICT CCIGACTACT GATGACCACT CACABACCA ATTSACCABA TIGIGATIGA AGATACTIAN ICCCCICCIC CCICIOGCIC CCCGICICAT TICTOTCCAC ICCATICICI CICCCICICI ANCIOCCITIC GITTICGIOTA GAITGACGCG TITICTITIGIA ATTICAGIGI TICTGACAAG INFORMATION FOR SEQ ID NO: 242: E Ě SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 242: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1146 base pairs 1661 360 240 480 420 300 180 120 1080 1020 60 540 720 660 960 900 780

30

CAAGGGGAGC COTGGAGCGC CACTCAGATT TOCCACCCCT GAACCCCACC TOGTGGTGGG

540

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90

CHOROSCETT TOTALOGOSE TECCHOSCIE TOTOGOSCIEE GOCATIECTO CTOCOTIECE

CAGATYTEAG GOCTETGGCA AGAAATCCTA GTTTCGTCCT GTCTTCCCTG GGCTTCACTG

GTCTCATCTT TGGACTCATC ACCTGCCTGA CCGGAGTCCT GGGTGTGGGC CTGGGTGTGG OCOTOGRACIO TOGOGRAGACO COACCOTOCO TROCCOGRAGA CICCIOCIOTI TOCICIORACA

CCCCOTTCCGC CACTCCAACC CCCGGGCTGA TCCCCTGGTC TGTGCCACTG

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TOSCCACTTA TATTTTCATC TTCATTOSAG AGACCCTCCT STCCATGAAC TGSSCCATCG GEORGETIGGG CICTIGEAGES TRICETOTISC TRICECOTIGG CIGGGESSIG GOTAGEATICG

GTCCCACCTG CTGGGTGATG CTGGGAGCCC CTACCTCATT GGCCTGATCT

TOTOCTOTAC GIGGIGATICS CTACCCGACG CTCCACCGCC GAGGCCTTCC

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960 900

1080

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CATETTEATT GAGGECGACE GEOGGEGE ACAGETGEAC GIOCAGGECE TOETGEACGA

1260 1200

GENEROCOCO TETTOTTOGOG CACTOGOCOG COCACTETICO TOGOCACCON GCGCCGGAAC TGGCCCCCCT CCTTCTTGTC CGAGTTCCGG GCTCTGCAGT

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KIGCCTIGGGC CCACCCCACG AAGGGCCTGG GCCTAACCCC TITGGCCTGGC CCAGCTTCCA

GTOCTCATCT GAGAGGCTGC CGCTCACCTA CCTGCACATC TGCCACAGCT

ACAGACGACC GGATTOTGGT GCCCCAGCGG GGCCGCTCCA CCCGCGTGCC

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AGATCAGCCG

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GACCAGOGA SOGGARGETC AGCATOTTOT ACTITICACAT TOCOGTOGGO-ÁGTGGTOTGG GICGOOGRAGO CCROTTARTIC CACCATOGOG COCACTOTICA TIGOCORCOT CTITIGIGGOC TEATSCITTEA TECCCOGAGA GEATTITETOG CTOCTECTICE TGACECOGGG CETGOTIGGG

> 360 300

420

OCTACATIVOC AGOCIVICAAA GIGAAGGAIA TOGCIVGAGA CIGGCACIVG GCIVIVGAGG

TOLACACCOOG TCTAGGAGTG GTGGCCGTTC TGCTGCTGTT CCTGGTAGTG CGGGAGCCGC

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OTOCTTOCOG ACATOGAGCA GITOTTOAAC ATOGGGGACA GIAGOTOTGG GOTOATOCAG

69

TACAATCOGA AGTATETEAT GTGCGGGGG ATTGCCTTET GGTCCCTGGT GACACTGGG ACCOTOTICA TCTCCAOTTA CATGOTOTTG GCACCTOTOT TIGGCTACCT GGGTGACAG

240 180 120 5

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SEQUENCE DESCRIPTION: SEQ ID NO: 241:

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INFORMATION FOR SEQ ID NO: 241: (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1661 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

CCTCTGATCT TOCACCCCGT CTTCACCCCA GGGCTCCTGA A GIAACGGAAT AAAATTIGTA GCCAGACCCC AGGIGCCIGC ICICGICTIT CICIGGGIGG

GAGGGACCCT GGGCCGTGTG CCAGCTCCCA GACACTACMT GGGTAGCTCA GGGGAGGAGG TOGGGGTCCA GGAGGGGAT CCCTCTCCAC AGGGGNCACC CCAAGGGCTC GGIGCIATT 1620 1560 1500

468

WO 98/39448

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CARTE

(2) INFORMATION FOR SEQ ID NO: 243:

.) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1350 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDRESS: double
(D) TOPOLOGY: linear 3

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SEQUENCE DESCRIPTION: SEQ ID NO: 243: ¥

1260 1320 0801 1140 700 960 1020 120 8 240 8 90 20 480 540 909 8 22 780 840 8 GCTGTGCCAC GGCTGTTGCT TCGGTTATTT AAATAAAAG AAAGTGGAAC TCGAAAAAA TOTOTOGGAC TTGGAGAGAC ATCACTAACT GATGGCTCCT CGGTAGTGCT AGTECECCCA TCAGGCCACA CTGCTGCCAC CTCTCACACG CCCCAACCCA GCTTCCCTCT CCCAATOCTA TOSCCATGAC TOCTGAACCT GACAGGOGTG TGGGGAGTTC ACTGTGACCT TECTECOSECA GECOTOGAG GECAGAGESC CECTGAGAGECS CAGTTGCAAA CANGGETCAG AGCAGAGGG GEGGAAACCC GTTCGCCGAG CCCAGCGAGC TTGACAACCC TOCOCTOCCT CENCENTINE ACTOCCIOCT CETITIGICIG CTGGTACCGC CECATGTATA AGGETITICCG GAGIGACAGI ICAITCAAIT ICITCGIIITI CITCITCAITI ITCITCGICC AGGAIGIGCI CITHOLOGIC CAGGCCATTG GIATCCCAGG ITGGGGATTC AGTGGCTGGA TCTCTGCTCT GOTGOTGCCG AAGGCAACAC AGCAGTATCC GTGCTCATGC TGCTGGTCGC CCTGCTCTTC ACTESCATTS CTGTGCTAGS AATTGTCATG CTGAAACGGA TCCACTCCTT ATACCGCCGC ACAGGIGCCA GCITITCAGAA GGCCCAGCAA GAATITIGCIG CIGGIGICITI CICCAACCCI GOOGTECGAA CCGCARCTTG CCAATGCAGC CGCTGGGGCT GCTGAAAATG CCTTCCGGGC CTGCTACTTG AGGGAGCTGA CTTFAGCTCCC CCACCAGCCT ATGAGCCTCC AGCCCTGCC CCATTGCCTC CACCCTCAGC CACCCTICGA GAAAGCTCAG CCCACAGAA CCTAAGAACT ATGGCTCATA OCCITCAGCTG CAGCAGCCAC AGCTGAGCTG CTGAAGAAAC AGGAGGAGCT CTGCCCTGGG ROSCACAGOT ACTOSACAGA ACAATIOSCO COCTOTACOT TOTTITIOTO CAGITICAGOS CIGIATOCAC GOCCADOTTIC TOTGTGGAAA CCAACAATGG GGCAGGCTTT GGGCTTTCTA TCCTCTGGGT CAACCOGAAG GCAGAGGAGT TOGACCGAAG GAGNCGAGAG CTGCAGCATG CATGTACTAC CTCTGGATGT GCAGCACGST GGCTCTTCTC CTGAACTTCC CTCCTTTTTC CAGGACATCT CCATGGAGAT CCCCCAAGAA TTTCAGAAGA CCCGTGACCC CTGACTGGGA TGCCCTGGCC STCCCTANGG AACCCACGGC CTTTCAGCCA TCCCTCCTTG CAGCACTCAG 6 45 တ္တ 55 8 33 2 25 2

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PCT/US98/04493

INFORMATION FOR SEQ ID NO: 244: (2)

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SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1529 base pairs

) TYPE: nucleic acid) STRANDEDNESS: double) TOPOLOGY: linear 3 2 2 3

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SEQUENCE DESCRIPTION: SEQ ID NO: 244: œ. 8 120 180 240

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360 420

TCCCAGAGGC COGGGGGTTC CAGCTCTGCC TGTAGCAGAG CCCTGAGGAG GAGGAAGA AGTGATGGCT TATGGCGGCC CACTCACTAT GGTGGGCTGA GTGGAAGGTC CTTAACCATG TOCTAGOTIT CATTITICCA TITICSCAAT CACCTICAA GICTICOCCCC AIGACAGCAA AGAIGIGCI GAAAIACGIC CGGGAGAICI TITICAGCIA GGGCAIAAAC IGIGCACIGA ACTICTOCC GAGAGCAGCT GGAGGACAGC TGAGCTTCCA CTGGTGCTGC TGGGCCGACC GOCTIONGSCA ANGOGOCIUT CHONGCIUCT ACCIMINATGU CITUTAGGO CHGGCAGAIT CACCTCAGGC CAGAAGCCCC TGGACACTCC GGGCCTTTGGG GTGCCGTTCT GAGTGTGCGG AAGGCAGGAC TCAAAATGAG ATCCCATTTG ACTCCCTCTG TATGTACTGT GCCCTCTC GOCICTICAG GCICTGGAGT CCCAATTGTC TGTGTTAGTC AGTGACCAGG TTCCAGGGAÀ TGCAAAGAAA TCTCCTGAAG CCCACCTCTC GCAGGTTGAG GAGCCCAGAA THYCAGIGIG OCTITICGACA GGAATATATG AATAAATCAC TGCCATACAG GTTTTCCAAT ACACAAGTGC AAGTTCTGGG GIGCGCAACC GICCAGICCT GITCACAGCG CIGICIGOCI ICAGOICCAG GGAGCITIG AAGCAGICAA GCCITGICIT ICIACCCCAI TACCTIGCT TECECAGAGA CACTGAGGTG CTEGETETT TAATGTECTE GTTTGTTGCE GTAAGTTIÇT GIGTOCTION TITICITICAGI CACTCAGAGA TCACTCCTGG ACCTCTGGGG TIGGAGTITC TAGAAAITAC ACACAATTCC CCAATGCGTA AGTTGTGCTA ATGTCTTTCC AGGATTIGGA GICCICCAGG GICTCAICAI GGGAGIGAIT IGICAGGGGA AATRATGTCA TOTOGTGGTC CAACTTACTG GAACCAAAGA GACAGTACTT CCCCATCATC GAGGAAGCCT GITICAAAA TAGITICCAT CAIGAGICTA ICAAIGAGCT CAGCCAGCCT AGAAAGCAAA CGAGCTGCCC ACAGTTCTCT GCCCTGTCTG GCCACAGTOT ATAGACTOGT AAGCCAGACA GGCCTCCTCC CGCAAGCTGC TCACCTGTAC CTTGGTCCCC GGGCAGCTAG CTATAAAGCA AGAGGGACAG GAGACACTGA GGACAAGAGA TCACACCAGA GTACATGTCT CTGCCTCTGT ACAGTTTAGT AGGATCACTG CCAGGTGCAC TGGAATTGCT ITCCCAACTC CACCITCCCA CCCITTICITIT

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480 540 1140 1200 1260 1320 1380

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900 960 1020 801

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CAGGAATTGG TCACTCCCTC CCCACCCTCC TGAAGCTAGA GGAAGATTTG CTCAGATCCA TTAATTAAAG TOTGACAATG AGCTGCATGG TTTAGGGAGT CTTTGGGAGC CTTGGAAGTC 471 1500 1440

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 1537 base pairs

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25 AGCANTOSCT AAGCCCCAGG TOGTTGTAGC TCCTGTATTA ATGTCTAAGC TGTCTGTGAA TOCCCCTGAA TITTACCCTT CAGGITAITC TICCAGITAC ACAGAAICCT AIGAGGAIGG

30 CCACCTICAC ACTITICANA CIGANATICA ACACTITICA GAGACCCIGA AIGCITOICI

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SEQUENCE DESCRIPTION: SEQ ID NO: 246:

TOCHOUNTT GOCCHOUNCE CSCCGCGGTG GCGGTTGCTA TCGCTTCGCA GAACCTACTC

180 240

360 300 120

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CCCANATTTC TCTTATATGG GAGCTCGCCT GTGTAATTAC CTGTCCCATC ATCTGACAAT 540

ACTTANAGAT CAACCTGCAA AAGGGGATGA AGTTACTCGA AAACGATTTC ATGCATTTGT

AAGAGCAGAT ATTCTTCAGG TIGGICTICG AGAATIGCIG AAIGCCCIGI TITCTAAICC

AGATGCTTGG AAGGAAAAAG GAAAGATGGA TATGGAAGAA ATTATTCAGA GAATTGAAAA

25

(2) INFORMATION FOR SEQ ID NO: 247:

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SEQUENCE CHARACTERISTICS:

COTTOTCCTA GATOCAAACT GCAGTAGAGA TOTAAAACAG ATOCTCTTGA AGCTTOTAGA

ACTOCOGOTOR AGUNACTOROS GONGAGUTOCA TOCARCUTOCA ACAUNTAGAS AAGCAACACO AGAAAATGAT CCTAACTACT TTATGAATGA ACCAACATTT TATACATCTG ATGGTGTTCC

1020

960 900 840 780 720

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1080 1140

TTTCACTGCA GCTGATCCAG ATTACCAAGA GAATACCAA GAATTACTTG AAAGAGAGGA GGANTCAGAG COTAAGCGAA AACAGTAAAG TTAAATTTTCA GCATATCAGT TTTATAAAGC GCATCATATT GATGATGAGA TOGACCCAGA GATAGAAGAA OCTTATGAAA AGTTTTGTTT CTITITITICCA GATTATIGAAG AAAATIGGAAC AGATTTATICC GGGGCTGGTG ATCCATACTT

<u>છ</u> INFORMATION FOR SEQ ID NO: 245:

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CTGAAGGACA AACAATCTTG TACTAAGAA

1529

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CARATTRAGG ATOTTGAGTT ATGITACTAR TOTATGCARC TITRATTTTTG TITRACACTA

AGTITIAGGTA TOGTGATITTA GCAGAACACA AGAGAGCAAG AAAATOTIGTC ACATCTATAC

TCTGCCAAAA TAAACTITAT TCCCTATAAC TTAAAATGTG TATATATATA

TTATGTACAG TTAKTTCTAC TOTTTTGGCT GCAATAAAAT CGATTTTGAA

ATAAAWRAAA

1537 1500 1440 1380 1320

(B) TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO:

GAGGCCCCTG AGAGCTCCAC CTAGTTCACA GGATAAAATC CCACAGCAGA ACTCGGAGTC

120

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(2) INFORMATION FOR SEQ ID NO: 246:

E

(A) LENOTH: 506 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear

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AAAAAAAAA AAGOGNOGCC GCTCTAGAGG ANCCAAG

240 180

TTGTGAGGAT TATCCTACTC TATCAGAATA TGTTCAGGAT TTTTTGAATC ATCTTACAGA 360 300

TACAACAGAT GATGCTTTGC AAGAACTTGT GGAACTCATC TATCAACAGG CCACATCTAT 480 420

TAGCCCACAG AGTGGCAACT TCCGCCAATT GCTACTTCAA AGATGTCGGA CTGAATATGA 600

660

8

AAAAAAAAA AAAAAAAATT TGGTGG

ATATTACTTT TTAGTTTGAT ACTAAGTATT AAACATATTT CTGKATTATT CCAAAAAAAA ATCCCAGCGG TCCTTACCAG AAAAAGCCTG TGCATGAAAA AAAAGAAGTT TTGTAATTTT TIGICACAGE AGIATGCIGT CTIGCCGACG GGGCCCTTAT TIACCGGAAG CTICIGITCA CTGTGTTGGC ACTGATACCA GAAACCACAA CATTGACAGT TGGTGGAGGG GTGTTTGCAC AGATOCTOCO OCTOGATATT ATCAACTCAC TOOTAACAAC AGTATTCATO CTCATCOTAT ATRACOTOCA GCCGARAATA AAACATCOCC CCTTCTCCTT CAGTGTGAAA GCCCACGTGA AGGCAGCCAG CTGAGAAGAG TTGAGGGAAAA GTGCTGCTGC TGGGTCTGCA GACGCGATGG

908 480 420

ACTOTYTOTO GGAGAACTTY AUCTYAACOT GGAGATCAAG GGAACAAAIG GACAGGTIAC

TANGGANGAC AATTIAAITT GIGCAGIAAA AITGIINAAG INGACAGGAT CAGITINGGA

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1260

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GCCCCTOTGA GGGCCAGCTC TGGAAAAACC TGGGAGTTGA TGCCGGAGGY TGGGAAGAAC ATBACECETT GRAACTOTOC CGAGTTECTT AAATETCAGE TGGGATECTG GACETGGGAG GICTITICTIT INCIGITITIG AGIIGGIGAG IGAGIGAAIA GGGIAACATG GGCCITCAGG

180 120

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SEQUENCE DESCRIPTION: SEQ ID NO: 247:

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear LENOTH: 1348 base pairs

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WO 98/39448

472

	TOTOCTICANG GOCAGOSTICS COTTGENACAC TOGTNATITOT GOGGOTOGGA GOCAGAGGGG	240
v	CTCCGGCTTT CTCTGAAATG AACACTGCTC TTCAGCAGTT CAAGTACTTG TTCTCAAAAC	300
,	ATTITICIDAL TGATIGGIAG GITTICATAA GCATIGITIC TITAAGGCAT GGAAAGGGAA	360
	GAATGCTCAA GCAAGTCATG TTTGTTTTCA GTGGGATGGG CCGGCGTTCT CACTGCTGGG	420
2	GOCTTCCCCT TOCATGTGGC ACCTTTGTGC AGGGCCACCA GGCAGACTCT TCCCACCTTC	480
	TCCCACTGAA GCACCAAGGG GCTTGAAGCG TAATTTGGCT AATCAGAGGC ATTTTTTTG	540
ž	TECTAGNATE TITCACACIT GIECAACGGT CITATITITI TAAAAGTICT GITGCTIGTA	009
3 ,	TINACAGGAA ACTAGAGAGA AATAGTITCT GAAGCCAGTT TATTGTGAAG ATCCCCAAGG	099
	GGAGGTTCGG TAGAGAAAA TAGTAAGCTG GTTTAGAAAC TGACGAGGGC AAACAGCCAG	720
20	GACCONTIGG AGAGGAATIT GCCAAAGAIC TACCCTGAGA TAAGGCCTGT CCAGTGTCTT	780
	CACCACGTGA ATRACCAGGG CTCCAAAGTG TTTTTCTGCT TTGAAAAAA AAATTCCACA	840
ý	AGCTTTTANA GOTOCATTTA AGANTOCATG TGACTTTAGA ATGGAACTGC OSGCOCTGGC	006
3	AACTOTCACG TOTOCTAGAA GOTTCGATGC CTCTGGAATG CATGTGATAC TCATCTCCAT	096
	TITIGITICCI TCAITICCAIT TITICITITI TACCACATOT CICCOTOTICS GICCIOTA	1020
30	AGAACTOGGA CACCTTGGTT ITTGTGTTAG ATTGAGGTGG GCAGCTGCAA TCAGCTTCTT	1080
	TATATGCAAA TTAGGCACGA CCCATCTGTG GTTCCCTGGT TGGTGGCTAA TGAAGTGAGG	1140
35	GENGGGAGGG ATGTCACCCC ANAUTHOGC CCTCCCATTG GCTTTGGCCA GGCCAGACAC	1200
3	TICACATECT TTACATGGTF CIGTGTAATT TTAAAGTTTA TGTGTATAAA GCGAAGGTGT	1260
	TICTGTGAAA CTGTATATTT TGTAAATAAA TATATTGCTA CTTTGAGAWR AAAAAAAA	1320
40	AAAAACTCGA GGGGGCCCG GTACCCAA	1348

45 (2) INFORMATION FOR SEQ ID NO: 248:

(i) SEQUENCE CHARACTERISTICS:

1500

1560 1620 1680 1740 1766

(A) LENGTH: 1766 base pairs
(B) TYPS: muclaic acid
(C) STRANDECNESS: double
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 248:

09	120	180
TTGCCGAATC GGCAGAGCGG CACGAGCGGC CACGAGAGCA GGCGGAGTAA AGGGACTTGA	SCHACCAGT 100006ATTA TTCTATTTCC CCTCCCTCTC TCCCCCCCG TATCTCTTTT	ACCOTICTE CCACCTEGE TEGGETASCA TGGEGGAGG TEGGEGGCA CTCAGTECCA
эс сассаванска во	oc cerecerere re	ся тоссоваесь то
co caccadose	TA TICIATING	GC TCGCGTASC
TC GCCAGAGO	от тоссоват	TC CCACCCTC
OTGCCGAN	GOGAGOCA	CACCCTTC
55		9

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PCT/US98/04493

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360

600 660 720 720 720 900 900 1020 1140 1140 1130 1130 1140

TICCATCTICC TOSTOGICCT TOGRACCGA GCCGTCCGCG CCCGGCGGG GCGGAGCCC TORGRACICG TICCCGARIG TOCTITICCIC CCICICCCCT GCCCACCICA AGITTARIA CICTOTCACC ATCACCTICA GOATCTACAA GTCCGTCATC CAAGCTGTAC AGAAGTCACA CCATAATTAC ATGAATGCTG CCATGGTGCA CATCAACAGG GCCCTGAAAC TCATTATTCG CTGATTCCAA GAATGCCATC TGATAAAAA GAATAGAAAT GGAAAGTGGG ACTGAGAGG RICCICCOCCA ITIGGGAAAG IGGITICCIAC GICACIGGAC ACCGGITICIG AGCAITAGII ATAAGGITGT ACTITITCTIA CIATAAAATA AAAAAAAA AACTCGAGGG GGGCCCGGTA AGAGCOTICO COCGOCOTIGG GCACGAAGAG CTGCAGCTICO TOCTGTGCGG, TGCAGGATO CATTITICIGG ACAGATOTGA AGAAGACTGG GITTIGICITT GGCACCACGC TGATCATÓC AGAAGGCCAT CCATTCAAAG CCTACCTGGA CGTAGACATT ACTCTGTCCT CAGAAGCTT TCICITICIS GIAGAAGAIC ISSTIGACTS CITGAAGCTS SCISICITICA ISTGGCTCA GACCTATISTY OGFOCTISTY TYAACGGAAT CACCCTTICTA ATTICTTISCTIG AACTISCTICAL TITICAGIGIC CCGATIOICT AIGAGAAGIA CAAGACCCAG ATIGAICACT AIGIIGGCA CAAAAAAAG GCAGAATAAG TACATGGAAA CCAGAAATGC AACAGTTACT AAAACACCAT TIAATAGITA TAAGGICGIT ACTIGIACIA IGAAGGAAAA TACICAGIGI CAGCITIGAG recagnitie accaetagne macteager arceantata gritigecen taagaagiea TIGGAGCCCI CAANTCCIAT CITCCIGCCC CACAAIGIGA GCAGCIACCC CIGAIACIC AGTCAGCAGO CATOCTGCGG TGGCGGTCAC TCCCTCTGCC ACTATCCCCA GGGAAGGAA GOTTICCCIG GCAGOTTICA GIGICATCAG TGIGGITTICT TACCICATCC TGGOTCTIC COCCCOAGAT CAGACCAAGT CAATTGTTGA AAAGATCCAA GCAAAACTCC CTGGAATCG AAGAATCAAA TTCATAGGAT AAGTCAATAC CTTAATGGTG GTAGAGCCTT TACCTGTAG TICADAGGG AAGATIGGA GGIAAGAGAG AAAATGAAAG AACACCICTG GGICCTICT TITICITIAA IGAITIAACI AICAACIIGA IAAAIAACIT AIAGGIGAIA GIGAIAAITI CIGCATICCA AGCITITITI TIBATITIGGI GITITICICC AICCITICCC ITTAACCCI AGIATCAAGC ACAAAATTG ATGGACTGAT AAAAGAACTA TCTTAGAACT CAGAAGAAG TGATTAACTT ATGAAAAAT TATTTGGGGA CAGGAGTGTG ATACCTTCCT TGGTTTTTT CCCAAATCGC CGGATATGAT CGTAAA S 2 8 ဓ္က 35 8 45 15 23

(2) INFORMATION FOR SEQ ID NO: 249:

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(i) SEQUENCE CHARACTERISTICS:

TYPE: nucleic acid LENGTH: 2664 base pairs

STRANDEDNESS: double TOPOLOGY: linear

S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9092

5 20 2 8 35 ઝ 25 3 50 AAGAGCTGCA GCTCCTCCTG TGCGGTGCAC GATCTGATTT TCTGGAGAGA TGTGAAGAAG ассамиссот созсассова свясвясява мессамями сстоссоса сстоявимс TASCATORCO GARCOTORGO GRECARTORA TEREFICIA TETERIORIE GIECTIOGRA TYPOCCERCE CICIETCOCG OCCOGNATET CTYTTCACCC TRETCCCACC CICGCICGGO AGTOTOCTOG GAGCAGGOGG AGTAAAGGGA CTTGAGOGAG CCAGTTOCCG GATTATTOTA GENATICACCC TICTHATTICT TOCTGARCTO CICATTITICA GIGICCCGAT TGTCTATGAG GACTOCTTGA AGCTGGCTGT CTTCATGTGG CTGATGACCT ATGTTGGTGC TGTTTTTAAC ACTOGOTITIO TETTIGOCAE CACOCTGATE ATOCTOCTITI CECTOGOAGE TITICAGIGIE THOGRATUTE CHECKAREET TECCETTIAA CECTEAGUAT CAAGCACAAA AATTGATGGA TACTATGAAG GAAAATACTC AGTGTCAGCT TGAGCCTGCA TTCCAAGCTT TTTTTTAAT CTTCHARAGA TCCARGCARA ACTCCCTGGA ATCGCCRARA ARARGGCAGA ATRAGTRCAT AAGTACAAGA CCCAGATTGA TCACTATGIT GOCATCGCCC GAGATCAGAC CAAGTCAATT TACAAGTEEG TEATECAAGE CTGATRARAG AACTRICITA GRACICAGAA GARGRARGAA TCRARITCAT AGGRIRAGIC GGAAACCAGA AATGCAACAG TTACTAANAC ACCATTTAAT AGTTATAACG TCGTTACTTG GOGGACAGGA GIGIGATACC TICCTIGGTI TITITITIGCA GCCCICAAAT CCIATCTICC CAGCTATICCA TTATAGTITT GCCCTTAAGA AGICATGATT AACTTATGAA AAAATTATTT GAGAGAAAAT GAAAGAACAC CICIGGGICC TICIGICCAG TITICAGCAC TAGICTIACI AATACCTTAA TOOTGOTAGA GCCTTTACCT GTAOCTTGAA AGOOGAAAGA TTGGAGGTAA TOCCCCACAA TOTGAGCAGC TACCCCTGAT ACTCCTTTTC TITAATGATT TAACTATCAA TITICITACCI CAICCIGGCI CITICICICIG TCACCATCAG CITICAGATC ACAGGGCCCT GAAACTCATT ATTCGTCTCT TICTGGTAGA AGATCTGGTT ACATTACTOT GICCICAGAA GCITICCATA ATTACATGAA IGCIGCCAIG TGTACAGAAG TCAGAAGAAG GCCATCCATT CAAAGCCTAC 1380 1320 1260 1200 1140 1080 1020 660 600 540 480 420 360 90 240 180 120 960 90 840 780 720 8

> ઝ မ 25 20 7 5 S GTITTIMACTG GTTTCATGTC CTAGTAGGAA GTGCATTCTC CATCCTCATC CTCTGCCCTC CGATTCCCAT TIGGGGGCAA GITTTITICT TCACCTICAA TAIGAGAATI CAGCGAACII CATGIGGGCT CCICAGITAT TGAGITITIG TGAICCTAIC TCAGICIGGG GGGGAACAIT CONCCANAGO ICITITOTANA GOGGICCCCC AMANTICCATT CANCGGGCCG ICGGITTITAN TOTTOGTICO TOTAAAATIT CAAATIGAIG TOOTATTAAT AAAAAAAAAA AAAACAMAAA CAGCTOCAAA AAGTAGTOGA AGGAAATTOT CTACOTOTCT TOGAAAAATT AGTTAGGAAT GGAATATGCA CTOOCGAGTT TAMAGTANGG GCTATGATAT TTGATGOTCC CAAAGTACOG GAAAGAAAAA TCATCTOTGA OTTCCTTCAG OTTCTCACTC ATAOTCATGA TCCTTCAGAG CTCAAGAGGT GAAATACAGA AAGCCTTTTT TICIIGATCT TIICCCGAGA TICAAAICTC AAATAAATOT CTOTAACIGC TOTGCACIGC TGTAAACTTG TTAGAGAAAA AAATAACCTG TODICCION COCCIGOCCA CONCAAGITI AATAAATAAG GIIGIACITI TOTTACIATA AAAGGTCGTG ANTGGGGGAA ANCC AAAAAAAAAA AAAAGGGCGG CCGCTCTAGA GGATCCAAGC TTACGTACGC GTGCATGCGA ACTGTGTGTA CGTGTCTGTG CGTGCAACAT AAAAATACAG TAGCACCTAA GGAGCTTGAA TOCCACACAG GATTITITIT TITTITAAGA AAAACCTATA GATGAAAAAT TACTAATGAA TGCCACATCC AGTICTITIC TITTGTTGCT GCTGTGTTTA GATAATTGAA GAGATCTTTG CCAGGAAGIC AGIGATIOIC TITTIOGOCT TCCCCTCCAA AGGACCTICT GCAGIGGAAG TIGOATOGGT AAAAGGIACC CTIGCCTTAC TCCATCTTAT TITCTTAGCC CCCTTIGAGI 2640 2580 2520 2460 2400 2340 2280 2100 2040 1980 1920 2220 2160 1860 1800 1740 2664 1680

(2) INFORMATION FOR SEQ ID NO: 250:

6

Ξ SEQUENCE CHARACTERISTICS: 8 (A) LENGTH: 865 base pairs TYPE: nucleic acid STRANDEDNESS: double

8

SEQUENCE DESCRIPTION: SEQ ID NO: 250: TOPOLOGY: linear

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8 S 8 TAGGAGGCCT GATTAARATG GTCCATCTAC TGGTCTTGTC AGGTGCCTGG GGCATGCAAA TIGGCOTTIC CICGGGGCIT TGGIGGGAIC GGIGICCICA GGAIGAGAIT TAGGGITTICC GOOTICCTICCO GATTIGAGGTC CCOGTTCCTA ACGOTGGGAT CGGTGTCCTC GOGATGAGAT COTOGGAGTG AGGTACCAGA TICAGCCCAT TIGGCCCCGA CGCCTCTKTT CICGGAATCC ICGGGGCITT CGGGATCITC ACCTAATATC CGGACTGCAA GATGGAGGAA GGCGGGAACC 300 240 180 120 69

8

TOTACOTOAC TOGACACOG TICTGAGCAT TAGTITGAGA ACTOGITCCC GAATGIGCTI

GTCACTCCCT CTGCCACTAT CCCCAGGGAA GGAAARGCTC CGCCATTTGG

GAAAGTOGTT

1620 1560 1500

AAAAAGAATA GAAATGGAAA GTGGGACTGA GAGGGAGTCA GCAGGCATGC

S

CTTGATRART ARCTTRIAGG TGATRGTGAT ARTTCCTGAT TCCRAGARTG CCATCTGATR

1440

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	TOTOGOTICAC CITOGICICA GSCITOCIGO TITITOCGAAG COTICOCOCA CAIACCITOS	360
•	GACTAGTICA GAGGAAGTIC TTCCCCTTCT ACTTCCACAT CTCCATGGGC TGTGCCTTCA	420
2	TCAACTICTG CATCTTGGCT TCACAGCATG CTTGGGCTCA GCTCACATTC TGGGAGGCCA	480
	OCCADETITIA CETIGETOTIC CIGAGOETIA COCTOGOCAE TOTCAAGGE CCETIGOCTGG	540
9	AACTOCOCIAC CACAGCTOCC ATOTOGGCCC TOCAAACCOT GGAGAAGAG CGAGGCCTGG	. 009
3	OTOGGGAGGT ACCAGGCAGC CACCAGGGTC CCGATCCCTA CCGCCAGCTG CGAGAGAAGG	. 099
	ACCCUAGTA CAGTOCICTIC COCCAGARTT TCTTCCGCTA CCATGGGCTG TCCTCTCTT	720
15	GCAATCTIGGG CTGCGTCCTG AGCAATGGGC TCTGTCTCGC TGGCCTTIGCC CTGGAAATAA	780
	GENECOTOTA GCATGEGCC TECATGCTAA TAAATGCTTC TICAGAAAA AAAAAAAAA	840
20	AAACTCGAGG GGGGCCCGGT ACCCA	965 20
25	(2) INFORMATION FOR SEQ ID NO: 251:	. 25
30	(1) SDQUENCE CHARACTERISTICS: (A) LENGTH: 2082 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPQLOGY: linear	30
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 251:	
35	TGGGGGGGN ANTGGGTGTC TGGCTCANGG ATTGCCNAAT CTGGAAATTC TCCATAACTT	95
3	GCTAGCTIGT TITITITIT ITTITIACA CCCCCCCGCC CCACCCCCG ACTIGCACAA	120
	TGTTCAATGA TCTCAGCAGA GTTCTTCATG TGAAAGGTTG ATCACCTTTG AAGCCTGCAT	180
40	CATTCACATA TITITICITIC TICITICCCT TCAGITCATG MCTGGTGIT CATTITICTGT	240 40
	STOTOTICTOT GITTATITT GITTGGAFTT THITTITIAA THINACTITT AGACCITECT	300
45	GIGTIGCOCA CCTTTTTICC AACCICCACC CTCACTCCTT CTCAACCCAT CTCTTCCGAG	360 45
}	ATGAAAGAAA AAAAAAGCA AAGTTTTTT TTCTTCTCCT GAGTTCTTCA TGTGAGATTG	420
	ACCTIGGAAA GGAAAAAAA AIGIGAAAIG TIAIAGACIT GCAGCGIGGC GAGIITCCAIC	480
20	GOGITITITI TITAGCAITIG TTATGCTAAA ATAGAGAAAA AAATGCTCAT GAACCTTCCA	540 50
	CANTCAAGCC TOCANCAACC TICTGGGTGT CACTTGTGAG TITTGGCCTT GTGATGCCAA	009
%	ATCTGAGAGT TRACTGGC ATTAAAAAA CTCATTCTCA TCTCATGCAT TATTATGCTT	\$5
3	GCTACTITOT CTTAGCAACA ATGAACTATA ACTGTTTCAA AGACTITATG GAAAAGAGAC	720
	attanatha taadaaaaa aagccigcat gctsgacats tanggiataa ttaittitic	780
9	CHITITITITITI CENTIFICATE ASSAURCE CENTICEARGE CHITACOLA CENTIFICA	09

900 960 1020 1080 1140 1200 1260 1320 1380 1440 1740 1500 1560 1620 1680 1800 1860 1920 1980 2040 2082 CITITOTITI AITOCCICAL GACTITITIG AOTITAGAAC AAACAGTOC AACOGTAGAG togggactaa aattictigcta ttgccgagaa gcagtctaaa atttattttt taaaaagağ GITAGGICIG TITAGCIGIA GAITITITAA ACGAITGAIG CACIAAAITIG ITIACIALTIG TAAGATATAA CCTGCAAGCA TATAATACAA AAAAAATTG CAAAACTGTT TAGAACGCTA TITTACAGIT GTATIGIGGI GCAGAACTGG AITHTCTGIA ACTIAAAAAA AAAICCACAG TITTAAAGGC AATAATCAGT AAATGTTATT TICAGGGACT GACATCCTGT CTTTAAAAAG GITCCIATGI GCITITCITI CATTITCAAT ICIGGITATA ICAAGAAAAG AATAATCIAC CCITICITICC ATGABATTIT GCATCTGCTC CAAAACTGCT TTGAGTTACT CAGAACTTC ACCTCCCAAT GCACTGAAGG CATTCCTTGT GCAAAGATAC CAGAATGGGT TACACATTT TGITAGCCCA GITAAAATGI ATGCTACAGA TAAAGGAATG ITATAGATAA ATTTGAAA TGATGTTAAG GGGGGTAGAG TTTGCAAGGG GACTGTTTAA AAAAAGTAGC TTATACAGGA ATTCABABAT GTATGTTTT TTTTCTTTTC ABABITABAG TATTTGGGAC TGABITTGCAC ACCTGGCAAA CATTGAAGAA CTCTTRATGT TTTCTTTTTA ATAAGAATGA GGCCCCACT AACTGCCCCA TTAITITITIGG THIGHTITIAT THITITITA TAITITITIGG CITTIGGIC TIGICAAAIG IGGAAIGCIC IGGGIITICIA GIAIAIAAIT IAAITICIAGI ITITIAIAAT TOTOCTITICA ACTIVAATAT AAGTIGGGTA TOTOTAGICT TIGCTATACC ACTGACTIGIA ATAAAATITA TGCAGTTATA AAAATGGCAT TACTGCACAG TITTAAGATG ATGCAGATIT AAATGAAAAG TAAATCTTAC CACAATAAAT ATAAAAAAT CTYGTCAGTT ACTTTTCTYT TIGAAAACCA AAGIAITIAAG AQQQQAAACG CCCCTGITITA TAICTGTAGG GGIAITITIAK TACATATITIT GCTGTGCAAA ATTGTTTTAT ATCTTGAGTT ACTAACTAAC CACGGGTGTT AATAATAAAC GGCATTTTTT TTTGAAAAA AAAAAAAAA AA S 2 15 2 23 ഉ 9 2

(2) INFORMATION FOR SEQ ID NO: 252:

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1482 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 252:

CAGCAGAGET GOCCCCGAGGG ACTIVETENCY GOCCCTGCCCCCCCCCCCCCCTT 60
GCCCCCCCCCCCCCCCCAGGG GCCCTAGGC GCCCTAGGAG CTGTTGTGAA CTTGTACCAA 120

479

GAGGTGATGA AGCACGCAGA TCCCCGGATC CAGGGCTACC CTCTGATGGG GTCCCCCTTG

180

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ANGGGICCC CCINGCIAAN GACCICCAIT CICCIGACCT ACGIGNACIT CGINCICICA OTGAACTIOT ACCAAGAGOT GATGAAGCAC GCAGATCCCC GGATCCAGGG CTACCCTCTG

180 120

45 6 ઝ 30 23 20 2 5 S GGAAACCAAA AAAAAAAAAAA AAAAACTCGA GGGGGCCCCG TA TOCCCTCATT GCTCTACAGG TTCTGGTATC CCAGTCATTA TICACCICAT CIGGATOTAT GGCACCATCT TCTTCATGCT GICCIGGICI GIGGCACAAC CCTACCTIIG GIGGAAAAAG CACAIGACAG CCATICAGCI GAICCAGITT ATAMACICTT TOCACCTOCT GIGGCACTOT COOTCIACAT IGIOTAIGAG TICOIGAIGT CGGGCIGGOT GAGCACOTAT AUGGCTAATC GGAAGCCCTT CCAGCTCCGT GGCTTCATGA TIGTCTACAA CTTCTCACTG CTAATGACCT CCATTCTCCT GACCTACGTG TACTTCGTTC TCTCACTTGG GCCTCGCATC CTOGONATAC AGCCTGTGGA GGCTGCTTAC TCAACTTGTG TCTTAATTAA AAGTGACAGA CCCCTCCCTG CCTTAAAACT TGGGAGAGGA GCACTCAGGG CTGGCCCCAC AAAGGGTCTC TOCTOCACOO CACACACTGA AGCAGTAGOT TOTOGGCCAA AGGTCAGGGT GGGCGGGGGC CACTGCACAT CTCCCAGTAC TACTTTATGT CCAGCTGTAA CTACCAGTAC ANGACGGGCA GOTGACCTTC CTACATOTCT TCCATCACTC TGTGCTTCCC TECTEACACA GAAGAGGTCA GEAATAATOT CACTOTOGAE CEAGTETEAC GETGTCTGCC ACTCCAGAGC TGGGGGCTAA AAGGGCTGTA CAGTTATTTC GACCACGGCT TITGGTICCTC ACCCACTICC CCCGGGCAGC TCCAGGGAIG AGGACTOCAC CYTAGGGCAG TOTOCGTCAG TOCCCTCTCC ACCTACACCT TROCCAAGGT ACTOTTATAC CAAGGGCIAG COGCTGCCCC GTGCACTTCA CCGTGCATGT CATAATGTAC CTGTACTACG GATTATCTGC CTTTGGCCCT TOTTCCTCTT CTCCAAGTTC ATTGAGCTGA TGGACACAGT GATCTTTATT GIGACCCIGI GGACTATICC AACAGCCCIG AGGCACTIAG GAIGGIICGG CITATOTOGT CAGGACTOAG CAGGGGACTG GCCCTCCCCT CCCCACAGCT CAAGGCCAAC TGAGAAGCAT GGCCTAGATA GGCGCCCACC GCAAAATIGGA GITTCTCCAAC

8 2 INFORMATION FOR SEQ ID NO: 253:

Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear Ξ LENGTH: 834 base pairs

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ž SEQUENCE DESCRIPTION: SEQ ID NO: 253

GGCACGAGCG CCGTTGCQCG CCTGGCCCCT ACGGAGTCCT TAGCCAGGAT GGAGGCTGTT

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1200 1140 1482 1440 1380 1320 1260 1080 1020 960 900 840 540 780 720 660 600 480 420 360 300 240

> 25 20 2 5 TODOTICADOS COTATACOTO OCOCTOTICAS COTOSAGENOT GOSCOTTIAGO GOSAGTOTOCO TACAACITET CACIGOIGGE ACTETECETE TACATIGIET ANGAGITECT GATGIEGGGE CCAAAGOTCA GOOTGGGCOG GOOCCTGOGA ATACAGCCTG TGGAGGCTGC TTACTCAACT AGGOCTIGGEE CEAEAAAGGG TETECTIGGEE TITTITECTEA CACAGAAGAG GTEAGCAATA CTAAAAGGGC TGTACAGTTA TTTCCCCCCTC CCTGCCTTAA AACTTGGGAG AGGAGCACTC TICCCCCOOO CAOCICCAOO GAIGIGOCCT CATIGCIGIC IGCCACICCA GAOCIGOOGO ACTORCETE ECCTECCEAE ARCTOSTETA CARROCAC OSCITIOSTI CETCACICAC TCAGTOCCCT CTCCAMCTAC ACCTOTGACC AAGOCTTATO TOGTCAGGAC TGAGCAGGG CTIGOGOCIO GCATCATOGO TAATOGGAAG COCTICOAGO TOCGIGOCII CATGATITOIO ANOTOACTOT GUACOCAGIC ICACICCICC ACCCCACACA CIGAAGCAGI AGCITCIGG

> > 540

480 420 360 岌

(2) INFORMATION FOR SEQ ID NO: 254:

834 780 720 660 600

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Ξ SEQUENCE CHARACTERISTICS: <u>0</u> (B) TYPE: nucleic acid (A) LENGTH: 1508 base pairs STRANDEDNESS: double

(D) TOPOLOGY: linear

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CCACCAACOT TOGGAGTOGA COTCATCAAT GAGOTTOTOG AGAACTTTOG CAGATOTOCO ATTICICCAT GCTCACCCTC CCAGGTCAGC GAGATGGTGA AGAAGCTGCA CGCGGCAACA AAGIGGICIG GICOGCAAGC CITIGICITT GICIGCCAGA CIGICATIGA GGAIGACIGC TIGAACITIT AAAATTITAG AICAGCAAAC ICTAAGAICC TAGAAIGGAA GCIGTICCIC Š SEQUENCE DESCRIPTION: SEQ ID NO: 254: 240 180 120 60

25

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GACTATTTCT TEGECTETE CAGCTECCAC CAGGAGGCTG TEGAGCAGAC CATCATEGET CTTCCCATGG ACCAGTITIGG TGTGCATCTC ATGCCGCATC TGCTAACCTT AGCAAATGAC GIGICTITICC TECTICCATG AGAGCCGAGG TICAGIGGGC ATTCGCCACG CATGIGACCI AAAATCTCCG AAGATGCCAT GAGCACAGCG TCCTCAACCT ACTAGAAGGC TTGAATCTCG AGGGTICCTA ACGIGCGAGI GCIGCTIGCA AAGACATIAA GACAAACICI ACIAGAAAAA ACCOTOACAG COATOTCAAG TATTITIGCAA GCATCCACCC TGCCAGTACC 600 420 360 300 540 480

PCT/US98/04493

GGGATAGCTT TCGGGGGAGG AGAGACCTTC CTCTCCTGCG GACTTCATTG CAGGTGCAAG ITICCTIACAC CCAATACCAG GGAITITCAAG AGTCAAGAGA AAGTACAGTA AACACTAITA

TAMATICCTIC TOTCTTCCTG ACTGAATGAA ACTTGAATTG GCAGAGCATT TTCCTTATGG ICTIATOTIG ACITIAAGGG GAAATAATIT CICAGAGGAT TATAATIGIC ACCGAAGCCI

AAGGGATGAG ATTCCCAGAG ACCTGCATTG CTTTCTCCTG GTTTTATTTA ACAATCGACA AATGAAATTC TTACAGCCTG AAGGCAGACG TOTGCCCAGA TGTGAAAGAG ACCTTCAGTA

TCAGCCCTAA CTCTTCTCTC CCAGGAAGGA CTTGCTGGGC TCTGTGGGCCA GCTGTCCAAC CCAGCCTOT GTOTGAATCG TTTGTGACGT GTGCAAATGG GAAAGGAGGG GTTTTTACAT

CTCCTAAAGG ACCTGATGCC AACACAAGTA GGATTGACTT AAACTCTTAA GGGCAGCATA

TIGCIGIACA CATITIACAGA AIGGITIGCIG AGIGICIGIG ICTIGATITIT ICAIGCIGGI CATGACCTGA AGGAAATTTA TTAGACGTAT AATGTATOTC TGGTGTTTTT AACTTGATCA

PCT/US98/04493

WO 98/39448

	AAGGGTIAAA GAACTGAAGG AAATCTGCCA TTCTCAGTGG ACAGGCAGGC ATGATGCTTT
'	TGAAATTITA GIGGAACTCC IGCAAGCACT TGTTITATGT TTAGATGGTA TAAATAGTGA
	CACAAATATT AGAIGGAATA ACTAIATAGC TGGCCGAGCA TTICTACTCT GCAGIGCAGT
	GICAGAITIT GAITICAITG ITACTATIGI TOTICITAAA AAIGICCIAI CITITACAAG
01	AGCCTTTGGG AAAAACCTCC AGGGGCAAAC CTCTGATGTC TTCTTTGGGG CCGGTAACTT
	GACTOCAGTA CTOCATTCAC TCAACGAAGT CATTOGAAAA TATTGAAGTT TATCATGAAT
7	TITIGGITTICA GGAAGCCACA AATTIGGCAA CCAAACTICA TAITICAAATG AAACTICCTG
	GGMAITTCCG CHGAGCTCAC CAGGGTAACT TGGAATCTCA GCTAACCTCT. GAGAGTTJACT
	ATHANGHANC CCTANGTGTC CCANCHGTGS AGCACATTAT TCAGGAACTT AAAGATATAT
20	TCTCAGACA GCACCTCAAA GCTCTTAAAT GCTTATCTCT GGTACCCTCA GTCATGGAC
	AACTCAAAIT CAATACSTCG GAGGAACACC ATGCTGACAT GTATAGAAGT GACTTACCCA
۶۲	ATCCTGACAC GCTGTCAGCT GAGCTTCATT GTTGGAGAAT CAAATGGAAA CACAGGGGGA
3	ANGATATAGA GCTTCCGTCC ACCATCTATG ANGCCCTCCA CCTGCCTGAC ATCAACTTTT
	TICCTAATIOT GTATOCATTG CTGAAGGTCC TOTOTATTCT TCCTOTGATG AAGGTTGAGA
30	ATGAGOGGTA TGAAAATGGA CGAAAGCGTC TTAAAGCATA TTTGAGAAC ACTTTGAGAG
	ACCAAAGGTC AAGTAACTTG GCTTTGCTTA ACATAAATTT TGATATAAA CACGACCTGG
3.6	ATTTAATOOT GOACACATAT ATTAAACTCT ATACAAGTAA GTCAGAGCTT CCTACAGATA
3	ATTCCGARAC TOTGGARART ACCTARGAGA CTITTARARA TAGGCTITCT TATAITTICAT
	ATTIGGAAGA AAAAGCCGTA AGTGTATGTA GACCACTTAA TCACTAAATA TCTTTGCCTA
40	TAGGACTCCA TTGAATACAT TAGCCATTGA TAATCTACCT GTTTAAATGG CCCCTGTTTG
	AACTOTONAG CITIGAAGAC CIACCIGITC ITOCAGAAGA GAACGITGAA AGTOCONNSI
45	ITCCITITIGE OTGATETETS TIGATGGEAE ICTGGAATTS TITCAGITAA GICATITIAG
}	ACATAGCATT TAITIAICACT GIGGAICTICT ACTIGITIGGG TOTTAIGAAT TCTTTGAAGA
	AATATTITT GAAGAGGTOT GOGAGGAAGG AATACATTIT ATAAAATGTT GTAGTGAAGC
. 20	CCACAAITGA CCITIGACTA AIAGGAGITT TAAGTAIGIT AAAAAICIAT ACIGGACAGI
	TACAAGAAAT TACCGGAGAA AAGCTTGTGA GCTCACCAAA CAAGGATTTC AGTGTAGATT
	TIGICITICI TGACTIAA GAACAAATG ACAAAGITIG AATGGAAAAG CCTGCTGTTG
3	TICCACAICT CGTIGCIGIT TACATICCIT TGIGGAGCT ACAICTICCT AAGCTITITA
	GCAGGIATAT GITGAACACT TCTGITTCAT GGITGAGACA GAATCAGAGG CCATGGATAC
09	TGACAACTGA ITIGICIGIT ITITTICICE GICTITITICC AIGACICITA IATACIÓCE

GIGCIGCAGT CITITAAICAT GCIGITITAAA CIGITOTGGC ACAAGTICIC TIGICCAAAT AAAATTTATT AATAAGATCT ATAGAGAGAG ATATATACAC TTTTGATTGT TTTCTAGATG TUTACCAATA AATGCAATTT GTGACCTGTA TTAAAAAAA NTAAAAAAAC TCGAGGGGG

IGATCAGCTIC TGAGGTGCAA CTTCTTCACA TACTGTACAT ACCTGTGAGC ACTCTTGGGA

CCCGGTAC

> TATGGGAGTA TCTGTTGCAT TAGGAACAAT TGAGGAAGTT TGTTCTTTTT TCCATCGATC ACCACAACTG CITTTAGAAC ITGACAAGGI AATTICIGIT CITTTICAGA ACAGIAAAGA

TATICTACACA CTCTGCTCTT CCTGTGCCTT AAATATGTGG TTGGCAAAAT CAGTACCTGT

GAGAGACTCA CACTTCTTTT CCATTATCAC TGACGATGTA GTCGACATAG CAGGGGAAGA GCACCTACCT GTGTTGGTGA GGTTTGTTGA TGAATCTCAT AACCTAAGAG AGGAATTTAT AGGCITCCTG CCTTATGAAG CCGATGCAGA AATTITTGSCT GTGAAATTTC ACACTATGAT AACTGAGAAG TGGGGATTAA ATATGGAGTA TTGTCGTGGC CAGGCTTACA TTGACTCTAG TGGATTTTCT TCCAAAATGA AAGTTGTTGC TTCTAGACTT TTAGAGAAAT ATCCCCAAGC

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 255:

(A) LENGTH: 2514 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 255:

8 3530 25 20 6 8 55 S 5 2 ARCCITIGAA GACCIACCIG TICTICCAGA AGAGAACGII GAAAGIGCCA IGITICCITI AACTOTOGAA AATACCTAAG AGACTTTTAA AAATAGGCTT TCTTATATTT GATATTTOGA TAAATAAAAT GIGAATIITIG TITAAAGCIT AGGCACATTA TITTITIGIGG GGTCAAAACA ATTTATAAGC AAAACCTGGA AAACCTACAA AATAAGTGTT GTGGTTTATC TAGAAAAATA TATOTTGAAC ACTICIOTIT CATGOTTGAG ACAGAATCAG AGGCCATGGA TACTGACAAC TCTCGTTGCT GTTTACATTC CTTTGTGGAG CCTACATCTT CCTAAGCTTT TTAGCAGGTA ANTIACCOGA GAAAAGCITG TOAGCTCACC AAACAAGGAT TTCAGTGTAG ATTEMPTATE ACTORGATE TETACTION GOSTOTIANG AATTEMPGA AGAMATATAT CCATTGAATA CATTAGCCAT TGATAATCTA CCTGTTTIAAA TGGCCCCTGT TTGAACTCTC CAGGAACCAG GATGTGGGTG CGAGGCGTGC TCCTGGCTGT TGCAGATTGC TGCACCCGGG TOGAMATAT TOCTOTTATT TETOSTGAAG AAAATCAATT TISTATASTT TATTICAATC TGATTTGTCT GITTTTTTTC TCTGTCTTTT TCCATGACTC TTATATACTG CCTCATCTTG TGACCITIGA CIANIAGGAG ITTIAAGIAT GITAAAAATC TATACIGGAC AGITACAAGA TITIGAAGAGG TGTGGGAGGA AGGAATACAT TITTATAAAAT GITGTAGTGA AGCCCACAAT TOCGTGATCT CTGTTGATGG CACTCTGGAA TTGTTTCAGT TAAGTCATTT TAGACATAGC AGAAAAAGCC GTAAGTGTAT GTAGACCACT TAATCACTAA ATATCTTTGC CTATAGGACT CATTTATTGG ACTITICISGI GCAMAAAGAT SITCAAGCCT TATITITATAC TISCCISCCC (2) INFORMATION FOR SEQ ID NO: 257: TICTIGIGIA AATICIC TOTTGAACIT AAAGAAACAA ATGACAAAGT TIGAATOGAA AAGOOTGOTG TIOTTOCACA (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: AGTGAGCTGC AGCTCTAAGA AGACCTGTTC TTTTGAATGG AGAGTAGCAT (D) TOPOLOGY: linear (C) STRANDEDNESS: double (B) TYPE: nucleic acid (A) LENGTH: 689 base pairs CITICICITY ATTITICICIT 2340 2160 2280 2220 2100 2040 1980 1920 1800 1740 1440 1860 1680 1620 1560 1500 180 120 60

GTACTGCATT CACTCAACGA AGTGANTGGA AAATATTGAA GTTTATCATG AATTTTOGTT CCCCAGAGCT CACCAGGGTA ACTIGGAATC TCAGCTAACC TCTCAGAGTT ACTATAAAGA TITIGATITICA TIGITACTAT IGITIGITICTI AAAAATGICC TAICITITAC AAGAGCCITT ATTAGATGGA ATAACTATAT AGCTGGCCGA GCATTTGTAC TCTGCAGTGC AGTGTCAGAT AAAGAACTGA AGGAAATCTG CCATTCTCAG TGGACAGGCA GGCATGATGC TTTTGAAATT CIGCTITIAG AACTIGACAA CGIAATIYCI GIICITITIC AGAACAGIAA AGAAAGGGGI GRATCIGITG AAGTIGGGGAT TAAATAIGGA GTATTIGTCGT GGCCAGGCTT ACATTIGTCTC CIGCCITATG AAGCCGATGC AGAAATTITG GCTGTGAAAT TICACACTAT GATAACTGAG 2 CAAAACATIC TIGIGIAAAT ICICITAAAC AITIGATAAA CAGCIICACA AITC AAAAATATGG AAAATATTGC TOTTATTTTT GGTGAAGAAA ATCAATTTTG TATAGTTTAT CATCTIGATT TATAAGCAAA ACCIGGAAAA CCTACAAAAT AAGIGITOIG GITTAICTAG ACAGCACCTC AAAGCTCTTA AATGCTTATC TCTGGTACCC TCAGTCATGG GACAACTCAA AACCCTAAGT GTCCCAACAG TGGAGCACAT TATTCAGGAA CTTAAAGATA TATTCTCAGA TGAGGAAGCC ACAAATTIGG CAACCAAACT TGATATICAA ATGAAACTCC CTGGGAAATT GOGAAAAACC TCCAGOOGCA AACCTCTGAT OTCTTCTTTG CGGCCGGTAG CTTGACTGCA TTAGIGGAAC TCCIGCAAGC ACTIGITITA IGITIAGAIG GIATAAATAG IGACACAAAT TCTTCCAAAA TGAAAGTTGT TGCTTCTAGA CTTTTAGAGA AATATCCCCA AGCTATCTAC TICANICIAA ATAAAAIGIG AATTITIGITT AAAGCITAGG CACAITATTI CACGCTGTCA GCTGAGCTTC ATTGTTGGAG AATCAAATGG AAACACAGGG GGAAAGATAT INFORMATION FOR SEQ ID NO: ž E CATTAGGAAC AATTGAGGAA GITTGTTCTT TITTTCCATCG ATCACCACAA CTTCCTGTGC CTTANATATG TGGTTGGCAA AATCAGTACC TGTTATGGGA SEQUENCE CHARACTERISTICS: TCGGAGGAAC ACCATGCTGA CATGTATAGA AGTGACTTAC CCAATCCTGA SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear 3 (B) TYPE: nucleic acid (A) LENGTH: 2357 base pairs STRANDEDNESS: double 256: TITGIGGGGT 2514 2400 600 540 480 420 360 300 240 660 180 120

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AGAGETTEEG TECACEATET ATGAAGEEET CEACETGEET GACATEAAGT TTTTTEETAA

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GICAAGTAAC TIGGCTTIGC TIAACATAAA TITIGATATA AAACACGACC TGGATTTAAT

GTATGAAAAT GGACGAAAGC GTCTTAAAGC ATATTTGAGG AACACTTTGA CAGACCAAAG TOTOTATISCA TITOCIGNAGG TCCTOTOTAT TCTTCCTOTO ATGAAGGTTG AGAATGAGCG

GOTGGACACA TATATTAAAC TCTATACAAG TAAGTCAGAG CTTCCTACAG ATAATTCCGA

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1320 1260 1200 PCT/US98/04493

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900 960 1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 1740 1800 1860 1920 1980 2040 2100

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2160

2220 2280 2340 2377

GTACCCCTCT TGTGTGGACG

GCTGTTTGGC AAAGACACAG TGAACACTAG TCTGAATGTA TACCGAAATA AAGATGCCTT GOSCITICCC TACGICCCAG AGCCCTATTA CCCGGAATCT GGATGGGACC GCCTCCGGA COGAGICITI GCIOCCGAAG CIGIGACIGC CGAITCGGAA GICCITGAGG AGCGICAGAA CATGGAGGTG CCGCCACCGG CACCGCGGAG CTTTCTCTGT GCACOTCCCC GAGGMYTTGA AGTCCTGAGC GCTCAAGTTT GTCCGTAGTC GAGAGAAGGC GCOTGOCTGG TGGCTGGTGG CATAATTGGA GCCTTGCTGG GCACTCCTGT AGGAGGCCTG AGAGGGTTAA CCTGGGTCAA ATGCACGGAT TCTCACCTCG TACAGTTACG CTCTCCCGG TORRAGISCT CIGARCITGA ARCICACIOG AGRGCIGARG GGRGCIGCCA TOICCGAIGA ATTGAACCAC TECTAAACCT TECTAGAAAC CETTEAGTAA TAGATAAACA AGACAAGGAC CTCCCTOAGA AAATTGAAAG TAGTTTACAG GAAGATGAAC CTGAGAATGA TGCTAAGAAA CTGATGGCAT TTCAGAAGTA CTCTGGTGAG ACTGTTCAGG AAAGAAAACA GAAGGATCGA GCARAGATT ARAGITGART TITRCROTTR ARABARARA ARABARARAR ARA ATATATOCAT ACATGAATAT ATCCACCCAC CTAGATTITA AGCAGTAAAT AAAACATTTC TOOTOGCACT GGCTTGCTCT TOTCTTTTTC TTTTCTTTTT AACTAAGAAT GGGGCTGTTG ATTOCCHACAG ACAGOCCACT CTTTGGTCAG CCTGCTGACA AATTTTAAGTG CTGGTACCTG AMOGCACTCC ATGAGCTAAA ACTGGAAGAG TGGAAAGGCA GACTACAAGT TACTGAGCAC AAGCCATTIT GTAAITIGCAG GAGCTGTCAC GGGAAGICIT GANAAACCCA AAGATGCAGA CAATCTCTTT GAACATGAAT TGGGGGCTCT CAATATGGCT (2) INFORMATION FOR SEQ ID NO: 260: Ê Œ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 260: ٤ N) LENGTH: 1262 base pairs

HTYPE: nucleic acid

STRANDEDNESS: double

HTYPOLOGY: linear TITAGGATAA ACGTAGGCCT AGAGCATTOT OCCUATITICS 1140 1193 1080 1020 960 840 780 720 660 600 540 480 420 360 300 240 180 900 8

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6 ઝ ၶ 25 8 5 5 GCATTACTAC GAAAAGAAGA AAGAGCAAGT CTTCTTAGTA ATCTTGGCCC ATGTTGTAAG GAATTITGAT CITTAGAGAC ACTAGITITIG GCCAACITAA GAITITACGI TAATTITTAG ANTARACTIT CACAGICCAG TATCCAACAG GAACTGIGIG IGICTIAAGA CCGAAGITCA CTTCGGTTAT TTAAGCGGAA GACTACTTGC CATGCTCCAG GACATGAAAA GACTGAAGAT ACCATAAAAC AAGCTTACAC GAGTSCTCCA ATGGTAGACA ATGAATTACT TCGATTGAGT OCCUTTOTICCT TCAGACCIGGA TICTICCAATT CGAAAGCAGC TICTIAAAAA IGAGAAGCOC ACAGACCACG TGAAAGGGAA TOCTGGTCTA GCTGGCGTGG TATGTTTATA GGCGAATTTC 3 CANCITECTI GOGATOTOGO CITITITIGGAA GGAAAAAAAT INCCCCAAAG GCAAATCCC CATTAGICAT CAACATTACA TOTTTCATGC TICAGATATI TIACIGCITG TOTCCTTATI GICTITITIT TAATTAATAT GIGIGCATIG TIACAAIGIA IGIIGGAIGI CITTIGACCC COCCTITIOT TAAAACIGAA GAITITIGGAA AAIGGITIGIC ACIGCICTIC CAGCCTAIGA ATAGTATITG ACACTCATGC AAAATAATGT GAAAACATCT AGATTTAGTA GITTATICIG GTTIGACAAG CATTAGTGAC AAAGGCAGAA AAGATTTATC AGCCATGCTA AAAGAGTGAA ATATOCIATT TITOCIACIG TETTECTICA GCAGIGCATA TICITITIGCA AAGITETITIG GTTGGACAGC TTTAAACAGA GTTGATGGTA CTTCAAATAT AGCTCATTGA TACTTAAGGG CCCTTTCCTT GITTIAGAIT TACTTTGCTC TICGITAATC TIATTCCTGA TGATCTAGAA TAAANGCTIT TITIGITANC AGAGATIGIG TACTATITIT ATITITAATA AANGTANCTI AGCAGAAGGA AGCCAAAATA GTTTTTTCCT TTTGAAAGTT TTTTAAAAAT TATTTCATGG TITATICAAA AACAAIGIGI ICAICAAAGI AATIGCICAC AIIGIGCAGI ACTAIGIIGI ATATTTTTOT GAAATOGAAC CATOGATTTA TOTCTOGATC ATCCATACAG AACCAACAAT 1260 1200 1140 1080 1020 960 900 840 780 720 660 600 540 8 240 180

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ATTICCAAGG ACICCAAAGC GAGGCCGGG ACIGAAGGIG IGGGIGICGA GCCCICIGGC

TOTANTOGOE GIOGOCOCOGO COCTOGOCOTT TOCOCOGIOG GGCGGGACTT

cenerates

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 259:

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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2 INFORMATION FOR SEQ ID NO: 261:

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Ξ SEQUENCE CHARACTERISTICS: 3 LENGTH: 1179 base pairs

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 261:

55 8 CAACGGCTCG GCAGCCAGCC ATGTCCTGCA CCCAGGACAG CGGCCCTGGG CTACAAGGAC GOCAMACTITI COCCCAMNGC TICGAMACTI GCAMGCCGAM ACCITGAMIC GITAMAMGIT GOGITIOCONO GGOGOCOTOG COCGAAGAAG CGCAAITIGGO GITICOGCGAA CGITIGGCOCT 180 120 60

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CC TGTCAAACAA CAGTGGCAAA AATGTGGAAC TGAAATTTIĞT

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			CCGATGGAGT CTTATATCCC TOTCAAACAA CAGTGGCAAA AATGTGGAAC TGAAATTTIST
	CTGGACTICA TCTTCCTGCG CCGACTGCG CGGGAAGGG GAGTTTCAGA CTGTGAAGGA	240	GGATTCCCTC CGGAGGCAGT TTGAATTCAG TGTAGATTCT TTTCAAATCA AATTAGACTC
v 1	COTCOTOCTO GACTOCCTOT TOGACTICIT ACCCOAGGG GTGAACAAAG AGAAGATCAC	300 5	TCTTCIGCTC TTTTATGAAT GTTCAGAGAA CCCAATGACT GAGACATTTC ACCCCACAAT
•	ACCACTCACG CTCAAGGAAG CTTATGTGCA GAAAATGGTT AAAGTGTGCA ATGACTCTGA	360	ANTOGOGOAG ACCOTOTATG GCGATTTCCA CGAAGCCTTT GATCACCTTT GTAACAAGAT
	CCGATGGAGT CTTATATCCC TGTCAAACAA CAGTGGGAAA AATGTGGAAC TGAAATTTGT	420	CAMPSON ASSAULTA ASSAURT ACCORDING CONTINUED
01	GGATTCCCTC CGGAGGCAGT TTGAATTCAG TGTAGATTCT TTTCAAATCA AATTAGACTC	480 10	CTISSISAGG GGCTITIAGGC CCGCCTCTCA TGAAATCAAG ACCCTTCAAA GGTATATGTG
	TCTTCTGCTC TTTTATGAAT GTTCAGAGAA CCCAATGACT GAGACATTTC ACCCACAAT	540	TTCCAGGITT TICATCGACT TCTCAGACAT TGGAGAGCAG CAGAGAAAAC TGGAGTCCTA
5	ANTCOGGGAG AGCOTCTATO GCGATTTCCA GGAAGCCTTT GATCACCTTT GTAACAAGAT		TITICAGAAC CACTITICAGA COCCAAGIAT GAGIANCICA TGACCCTITCA
?	CATTOCCACC AGGAACCCAG AGGAAATCCG AGGGGGGCC CTGCTTAAGT ACTGCAACCT	099	TYZARTICZTA ANTORGACZA CACTETYCCCT CANTEGGACAT GAAAGAAGAC AGACTTTYAA
	CTIGOTICAGE GECTITIAGGE COGCCITCIGA IGAAAICAAG ACCCITCAAA GGIAIAITGIG	720	OCTURACIO OLI POLI DE CONSTRUIR DE CONTRACORA PARTICIONALE DE PARTICIONAL DE CONTRACTOR DE CONTRACTO
20	ITCCADGITT ITCATCGACT ICTCAGACAT IGGAGAGCAG CAGAGAAAAC IGGAGTCCTA	780 20	ATTACOATT TOACOCTA COATRIACT COCACOCTA TACATTACTACT TOACTACTACT
	TITICAGAAC CACTITICIGG GAITICGAAGA CCCCAAGTAT GAGTATCTCA TGACCCTTCA	940	CTACATTGCA CAGOTTCAGC CAGTATTCAC GTGCCAGGAA CAGACCTACT CCACTTGGGT
, 25	TOGACTICOTA ANTGACACA CACTOTCCCT CATGGCACAT GAAGGAGAC AGACTTTAAA	900 25	ACCCTOCAAT TAAGAATCAT TTAAAAATGT CCTGTGGGGA ACCCATTTCA GACAAGACAG
1	CCTTATCACC ATGCTGGCTA TCCGGGTGTT AGCTGACCAA AATGTCATTC CTAATGTGGC	096	GAGAGAAAA NAANGAAAG AG
	TAATOTCACT TOCTATTACC AGCCAGCCC CTRIGIAGCA GATOCCAACT TTAGCAATTA	1020	
30	CTACATTOCA CAGGITCAGC CAGTATTCAC GTGCCAGCAA CAGACCTACT CCACTTGGCT	1060 30	
	ACCCTGCAÁT TAAGAATCAT TTAAAAATGT CCTGTGGGGA AGCCATTTCA GACAAGACAG	1140	(2) INFORMATION FOR SEQ ID NO: 263:
35	GAGAGAAAA AAAAAAAA AAAAAAAAA AAAAAGAGC	35	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 715 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear
40	(2) INFORMATION FOR SEQ ID NO: 262:	40	
	(i) SEQUENCE CHARACTERISTICS:	•	COSOCTIOSOT ATTTOCCTOG CACCATOGOG CCCAAGOGCA AACTGGGCAC GAGAGGAAG
	(B) TYPE: nucleic acid		
45	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	45	AACCAGNIAT TIGAAGAA CAGAGACT CIGAAGTICT ALCIGCGGAT CATALIGGG
	. And the state incommendation incommendation 15!		GCCAATGCCA TITACTGCCT TGTGACGTTG GTCTTCTTTT ACTCATCTGC CTCATTTTGG
	(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 461:		OCCIDENTIOS COMICESCAN TAGNOTICACA GISTATICAGO COACCIACOA CICHATICAGO
20	GGCMACTIT CCCCCANNG TICGAAACIT GCAAGCCGAA ACCTIGAAIC GITAAAAGIT	60 50	TOGATOGCAC GAGCAGCSTT CTTCTGAGGA TGGGGCCCTG ATGGATGGTG GCACGAGCTC
	GOSTIGOGNO GROCOCOTIGO COCGINAGANO COCANTITOSO GITICOGOGIA COTTOGOCOT	120	AACATGGAGC AGGGCATGGC AGACCACCTT AAGGATGTGA TCCTACTGAC AGCCATCGTG
	CHACGGCTCG GCHGCCHGCC ATGTCCTGCA CCCHGGACAG CGGCCCTGGG CTACAAGGAC	180	CAGGIGCICA GCTGCTTCTC TCTCTATGTC 1667CCTTCT GCCTTCTGGC 1CCAGGCCGG
55	CTGGACCTCA TCTTCCTGCG CCGACCTGCG CGGGGAAGGG GAGTTTCAGA CTGTGAAGGA	240 55	occentrace rectionast gaangreens ascereings reachedaga cagnoscales
	COTOGRACIOS GACTGCCTOT TOGACTTCTT ACCCGAGGGG GTGAACAAAG AGAAGATCAC	300	CONCRORAGE ACCORDANCE GRANDED COOCERCING ACCORDAGE ACCORDAGE GATGANGCIG
9	ACCACTCAGG CTCAAGGAAG CTTATGTGCA GAAAATGGTT AAAGTGTGCA ATGACTCTGA	360	THATACCAT TOACATTOTG OCCACAGGCC ACTOSCCCTG GOTGGCTCTG TCAGGGTGCA

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 265:

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() LENGTH: 1638 base pairs
3) TYPE: nucleic acid
C) STRANDEDNESS: double
D) TOPOLOGY: linear

E SEQUENCE CHARACTERISTICS:

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2) INFORMATION FOR SEQ ID NO: 265: 50

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3 6 CTCCAGOGOC CAAAAGCAGT CTGAGGTATT GGGTATACTT ATACTCTATA GGGTCGTTGA CCGACAGGAG TOGCTCTGTC AGGGTGCACA GCCCCTCATG CCTGGAGCAA TGAGGGTCTA CGGCGCAGA TGAAGCGGTT ATAGCCATTG ACGATTTXGC SACINFIGCCAC

780 720

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TAAATAAATT TCCCAGTTAA AGATTATTGT GACTTCACTG TATATAAACA TATTTTTATA

CITTATIGAA AGGGGACACC IGTACATICI TCCATCRICA

AGACTGATTG GAATTCTTTC TOTTGAAAAG

CACACACAAT AAAGAACCCC CTGTAAAGAC AAATAAATGA

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GAGGAATIGT GCTTTCACCA

GAATTTICTAA GGATTTICTOG CTTAAATATC ACCTAGCCTC

1638 1620 1560

GAAGATAGCG TOCTOTGATT

CATATATITG CAGTATGAAC TATTGCCTCT

GGGACGITGI TITICACATOG

TACATTCAAC TCTGATCCCG

GGGCCTTAGG

1500 1440 1380 1320

CCCCTGGTTC ACTOCAGACA GTOGCACCCC AGCACCAGAG CACAATGAGA AACGGCAGCG 660 600

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TAGCACTTAC GTAAAACATT TCAAGCAGGA CCCTAAGATG 35

TAGAGCTTTT TAATAGCACT AACCAATGCC TITTTAGATG TATTTTTGAT GTATATATCT ATTGAACAAA AATGITTICCA CIGGCTTIIG CCIGIAAGAA AAAAAAATGIA CCCGAGCACA

ATTATICAAA AAATCAIGIT TATTITIGAGI CCTAGGACIT AAAATTAGIC TITIGTAATA

AAGCTGAGCT TTTGATGCCA GGTGCAATCT ACTGGAAATG

TGTTTCCCCC ACAGTTTTAA TAAGAACAGA TCAGGAATTC

1260 1200 1140 ဗ

CATCTTOTOG CGAGCTOCTO ATACAGAAGA GATOGATATT GAAATGGACA GTGGAGATGA GAAACAGGAA GAGCGAGTAC AGCAAGTACG CAAGAAATTG GAAGAAGCAC TGATGGCAGA TIGGCITAAC ACATCICAAC COCTCIGCAA AGCTTTTATT GTCACAGATG AAGACATCAG CACAMBETET GEGECAATEA CAGGGCAAGT CTCCGCTGCT GTGGAAAAAGA ACCCTGCTGT

840

780 720

AGCCTAAGAA TATGATCAGG TAACTTTCGA CCGACTTTCC CCAAGAGAAA ATTCCTAGAA

1020

960

900

1080

GGATGTGATC CTTCTGGCTC CAGGCCGGGC CCTTTACCTC CTGTGGGTGA ATGTGCTGGG 540

CTACTGACAG CCATCOTOCA GOTOCTCAGC TOCTTCTCTC TCTATGTCTG

35

GCAGGGCATG GCAGAGTGAG TGTCCCCCAC CGCCAGCCCA GGCACCTTAA

480 420 360

ACGAGCAGCG TICICICAGG ATGGGGCCCT GATGGATGGT GGCATGGACC

GCTCGATGGC TITAGICIGG CAGIGIAIGG GGCCAGCIAC CACICIAIGA

30

GOGGCCAATG CCATTTACTG CCTTGTCACG TIGGTCTTCT TOGGCCTGGT TGGCCTGGGC TITACICATC TOCCTCATIT

25 AAGAAGCAGA TATTTGAAGA GAACAGAGAG ACTCTGAAGT TCTACCTGCG GATCATACTG 300 240 180

AAGTOCATGA GCTGCCGATG TGGTGCTTAG TGATTGCGGT TTCGGTCGCT CTCCCGTGTT TECEGOGETO GETATTIBEE TEGEACEATO GEGECEAAGG GEAAAGTGGG CAEGAGAGG

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TAGTOCATCA GATGTAACAG AACAAATTAT AAAAACCATG GAACTACCCA AAGGTCTTCA CCCACAACGA ATGAATGAAC AGCCACGTCA GCTTTTCTOG GAGAAGAGGC TACAAGGACT AATTYTCAAA CAACCGGTAA CCAAAGTCAC AAATCATCCT AGTAATAAAG TGAAATCAGA

AGGAGITIGET CCAGGTAGCA ATGATGAGAC CCTTTTATCT GCTGTTGCCA GTGCTTTGCA

660

90 540 480 420

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GCCTCAGITIG GCAAGGTACC TGGGAAATAC TGTTGATCTC AGCAGTTTTIG ACTTCAGAAC

AAGTGCTGGC AAGAGCGATG TCTACTACTT CAGTCCAAGT GGTAAGAAGT TCAGAAGCAA

240 180 120

GGATTIGCCCG GCCCTCCCCC CCGGATGGAA GAAGGAAGGAA GTGATCCGAA AATCTGGGCT

OTCOGOGAGO GOOGOOYCOG OGYCCAGGOG ANCCCOGGINO ACOGIAGAGOG GGAAGAGGAT GOCIACIAGOS GOCOGIAGOS GIBOCOGIOS COCOCOCOGO CIGIGIAGODOT INCOCITITOCO

60

CARTCAAAAT AAGGGTAAAC CAGACTTGAA TACAACATTG CCAATTAGAC AAACAGCATC

TOGAMAGATG ATOCCTAGTA AATTACAGAA GAACAAACAG AGACTGCGAA ACGATCCTCT

360 300

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO:

(A) LENGTH: 783 base pairs

15

(i) SEQUENCE CHARACTERISTICS:

5 (2) INFORMATION FOR SEQ ID NO: 264:

TIGGGTATAC TIATACICTA TAGGGTCGTT GAATAAATGG CTTAGAATGT GAAAAAAAA CAGCCCCTCA TOCCTOGAGC AATGAGGGTC TAGTCCAGGG GCCAAAAGCA GTCTGAGGTA ааааааааа аттт

> 735 720 660

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(2) INFORMATION FOR SEQ ID NO: 266:

٠,	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1455 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 266:		•
	CSTUCOTACT GCCATGCAGG TACCGGGTCC GGAATTCCCA GGGTCGACCC ACGCGTCGGC	09	
	TCACTTGGCA AGGTACCTGG GAAATACTGT TGATTCTCAGC AGTTTTGACT TCAGAACTGG	120	
	AAAGATGATG CCTHGTAAAT TACAGAAGAA CAAACAGAGA CTCCGAAAGG ATCCTCTCAA	180	
	TCAMARTANG GGTANACCAG ACTTGANTAC AACATTGCCA ATTAGACAMA CAGCATCAAT	240	
	TITCAAACAA COGGTAACCA AAGTCACAAA TCATCCTAGT AATAAAGTGA AATCAGACCC	300	
	ACANGGAATG AATGAACAGC CAGGTCAGCT TTTCTGGGAG AAGAGGCTAC AAGGACTTAG	360	
	TOCATCHCAT GTAACAGAAC AAATTATAAA AACCATGGAA CTACCCAAAG GTCTTCAAGG	420	
	AGTIGGICCA GGIAGCAATG ATGAGACCCT TITAITCTGCT GTTGCCAGTG CTTTGCACAC	480	
	ANGCIPCIGOS CCAATCACAG GOCAAGITC CGCTGCTOTO GAAAAGAACC CTGCTGTTTG	540	
	GCTTAACACA TCTCAACCCC TCTGCAAAGC TTTTATTGTC ACAGATGAAG ACATCAGGAA	009	
	ACHGGANGAG CGAGTACAGC ANGTAGGCAA GAAATTGGAA GAAGCACACACACAT	099	
	CTTOTCOCCA GCTGCTCATA CAGAAGAGAT GGATATTGAA ATGCACAGTG GAGATGAAGC	720	
	CTAAGAATAT GATCAGGTAA CTTTCGACCG ACTTTCCCCA AGAGAAAATT CCTAGAAATT	780	
	GAACAAAAT GTITCCACTG GCTTTTGCCT GTAAGAAAA AAATGTACCC GAGCACATAG	840	
	ACCTITITIAA TAGCACTAAC CAAIGCCITT TTAGATGIAT TTTTGATGIA TAIATCIAIT	006	
	AFTICAAAAA TCATGTTTAT FTTGAGTCCT AGGACTTAAA ATTAGTCTTT TGTAATATCA	096	
	AGCAGGACCC TAAGATGAAG CTGAGCTTTT GATGCCAGGT GCAATCTACT GGAAATGTAG 10	1020	
	CACTTAGGTA AAACAITTGT FTCCCCACA GTTTAATAA GAACAGAITCA GGAATTCTAA 10	1080	
	ATABATTICC CAGTIBABGA TTATTGIGAC TICACIGIAT ATABACATAT TITTATACIT 11	1140	
	TATTGAAAGG GGACACCTGT ACATTCTTCC ATCRTCACTG TAAAGACAAA TAAATGATTA 12	1200	
	TATTCACAGA CTGATTGGAA TTCTTTCTGT TGAAAAGCAC ACACAATAAA GAACCCCTCG	1260	
	TTAGOCTICC TCTGATTTAC ATTCAACTCT GATCCCGGGG CCTTAGGTTT GACATGGGAG 13	1320	
	GTGGGAGGAA GATAGGGCAT ATATTTGCAG TATGAACTAT TGCCTCTGGG ACGTTGTGAG 13	1380	
	GAATTGTGCT TTCACCAGAA TTTCTAAGGA TTTCTGGCTT AAATATCACC TAGCCTGTGG 14	1440	
	PARTITITY TCCT 14	1455	

v	(2) INFORMATION FOR SEQ ID NO: 267:
n	(i) CONTENTS (ii)
	TYPE: nucleic
10	(C) STRANDEDNESS: double (D) TOPOLOGY: linear
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 267:
2	COCCTOCAGT ACCOGTOCOO ANTICCCOOO TOCACCCACO COTCCCTGAC CCAGGAGAA
C	CTGCCTGTCT ACATCAGCCT GGGCTGCAGC GCGCTGCCCC CGCGGGGCCG GCAGCTGAA
	TATORICICT TCAGGGGGG CACCOTGTG CATTCATCTT TGTACCCCCA GCATCTAGC
20	STOTTOSCAT STAGTAGSCA CTCAAGAAAT STOTGTTGAA TGAAGGATGC CTGTGACAA
	CAAGOGAACT TTAITCTITC CTGACCCTTG CTCCTAIGAC ACACCTCCTC CTGACTGCC
Ÿ	CTGTCACTCC TTCAGAGCAG AACTCCTCTA GGGAACCTGG ATGGGAAACA GCCATGGCC
3	AGGACATECT GOSTGAAGCA GOGCTACACT TTCATCGAACT GAACAAGCTG AGGSTGTTG
	ACCEMBAGGT TACCEMBEAG ACCATAGAGE TCAMGGAAGA GTGCAAAGAC TTTGTGGAC
30	AAATTGGCCA GTITCAGAAA ATAGTTGGTG GTITTAATTGA GCTTGTTGAT CAACTTGCA
	AAGAAGCAGA AAATGAAAAG ATGAAGGCCA TCGGTGCTCG GAACTTGCTC AAATCTATA
y.	CAAAGCAGAG AGAAGCTCAA CAGCAGCAAC TTCAAGCCCT AATAGCAGAA AAGAAAATG
e.	acctrgrad ctatccoctt gratatgrac ctttctgtaa actrgragge gracaaaat
	AATTTATTGA CCAATTTATT TITCAGAAAT GAACTGAAAA TITCGCTTTT ARAGTAGGA
40	GGCAAAACAA AAAAAAGCCT CTCAAAAACCA AAAAAACCTC TGTAGCATTC CAGGGGCTT
	ACCANTGACC TRIFFICACAA GAGGIGGGGT GTAAGGAATG CAGCCCCTG AAGACAGCA
34	TACAAGTETE GGGAGCCAG TTTTAACATC AGTGCACAGC TGCTGCTGGT GGCCCTGCA
}	TOTACOTICT CACCICITAT GCTTAGTTGG AACTAAGCAG TITGTAAACT ITCATCCTT
	TITITIGIAAA TICACAAAGC TITIGGAAGGA GARGCAATAA AITITITIGKIT TCINAANIGG
50	TTGATG
55	(2) INFORMATION FOR SEQ ID NO: 268:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1003 base pairs
(B) TYPE: nucleic acid
(C) STRANDERNESS: double

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SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear

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20 2 5 8 35 30 25 Ś AAAAGCAAGG CTCCTAACTC ATTGGGAACA ACTGGACTAT GGAGTACAGT TTACATCTTC GOCACGOGAG CAGCCGOGCT GOTCCTGCTG CGAGCCGGGG GCCCGGAGTG GGGCGGCGA GTATGATICA ACTICACTICA TECTAMACAE AGETTETETE CIGAGIGIAE TAATTEECAA AGCATTCCCT TCTTCAGTGT TCCTGTTGCT TGGACTTTAA CAAATATTAT ACATAATCTG TCAGCTGTGG TTATTTAAAG CAGACTTACA TGTAAACCGG AATCCTCTCT GOGCATCACA GTGAGGGTGT AGTAGATAAA TTCAAGGAAA TAAGAGATTT GTAAGAAACT GOTACATTIC TGAAGAGGG CITTATAAGC AGGCIGGGCA GGCCCAGCIT ATAAGTIAAA CACTOTITCT CTCATAATOT GAAATGAGAA GTATTTACAT TOGAGGOCCA ATGCCTGGTC AATGCCACAA CTACATGGTG TTCGGATCTT TGGAATTAAT AAGTATTGAA ATGTTTTGAA GOGATOTACO TATTITIGCA TOCAGTGAAA GGAACACCTF TCGAAACTCC TGACCAGGOT AGCCGGGGTA TGTGGCTGAC ATATGCATTG GGAGTTGGCT TGCTTCATAT TGTCTTACTC GCAAACATGA ACGITGGAGI TGCCCACAGI GAAGIGAATC CAAATACCCG IGTCAIGAAC 2 ATTAAAGATT ATTITITAITA CCGTAAAAAA AAAAAAAAAA AAA AGGACCAGCT TAACTTATAA TGAATGGGCA TTGTGTTAAG AAAAGAACAT TTCCAGTCAT CTTCAAGTGC actgaaaaa aattitacag ctactgaatt tcttataagg aaggagtggt tagtaaactg ACGGAAGTTT TICACAATIT CICCAATAAT ICIATATITI CIGGCAAGIT ICIATACGAA INFORMATION FOR SEQ ID NO: 269: TOTTTTCAAG TOCAGATTTC CATTAAATGA TOCCTCTOTT TAATACACCT ATACAAGTTI 1003 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 8

E SEQUENCE CHARACTERISTICS: LENGTH: 1234 base pairs

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€ 0 TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear

55 50 8 ACATATOCAT TOSSAGTIOS CITOCITCAT ATTOICTIAC TCASCATICC CITCITCAST GTTGCCCACA GTGAAGTGAA TCCAAATACC CGTGTCATGA ACAGCCGGGG TATGTGGCTG CCAGTIGIAT CAGIGITGAT TCATTICATT ACTICCTACA GAGCAAACAT GAACGITGGA ATCAGCATCT ACAAGTAGCA TATTTTIGGAT GGTGTTTTGTG TGCTACTTCA AAGTAACTAG Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 269:

300 240 180 120

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35 5 INFORMATION FOR SEQ ID NO: 270:

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CASTICCCAC TRAIGAGGGT ACTITITITIGG TITTICCTICG CITAATATIG TGTATTOGIC

AATGAGGCCA TITTITACANT TATTAACGIT ACAG

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GCAGACTTAC ATGTAAACCG GAATCCTCTC TATACAAGTT TATTAAAGAT TATTTTTATT

TITICKCITGT TITATGIAAG YGGAIGIATA ICCICITGIT

TTATACAAGC

1200 1140 1080 1020

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TAGTAGATAA ATICAAGGAA ATAAGAGATT TOTAAGAAAC TAGGACCAGC TTAACTTATA

960 900 840

ATTOTOTTAA GAAAAGAACA TITICCAGTCA TICAGCIGIG GITATITIAAA

20

GCTTTATAAG CARGCTOGGC

AGGCCCAGCT TATAAGTTAA AGGGCATCAC AGTGAGGGTG

GIGCAGATIT CCATTAAAIG AIGCCICIGI TIAATACACC

TOGTACATTT

CTGAAGAGGG

TOAAATGAGA AGTATITACA TIGGAGGGCC AATGGCIGGT CCIICAAGIG CIGITIIGAA

GCTACTGAAT TICTIATAAG GAAGGAGIGG TIAGTAAACT GCACTGTIIC

TSTGATAATO

720 660 8 540 480 420 360

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GTICOGAICT TIGGAATIAA TAAGTATIGA AAIGTTITGA AACIGAAAAA AAATTITACA ATCCTAAACA CAGCTTCTCT CCTGAGTGTA CTAATTCCCA AAATGCCACA ACTACATGGT TCTCCAATAA TICTATATIT TCTGGCAAGT TICTATACGA AGTATGATCC AACTCACTIC CATTOGGAAC AACTOGACTA TOGAGTACAG TITTACATCTT CACOGAAGTT TITTCACAATT CATOCAGTGA AAGGAACACC TTTCGAAAACT CCTGACCAGG GTAAAGCAAG GCTCCTAACT GTICCIOTIG CTIGGACITI AACAAATAIT ATACATAAIC IGGGGAIGIA CGIATITITIO

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8 3 Ξ ž SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 270: <u>0</u> (B) TYPE: nucleic acid (A) LENGTH: 574 base pairs (D) TOPOLOGY: linear STRANDEDNESS: double

CACTGCTACC TGGTACTGCT TTCAGTGTGT TCCCCCTCAG CCCTCCGGCG TGTCAGGCAT AAAACCTCAG TCTGCCTGTA AATTTCAGCA AGCCGTGTTA GATGGGGAGC GTGGAACGTC TATOGATECE CATGAAGEEE TACTACACEA AAGTTTACEA GGAGATTTOG ATAGGAATOG ACTOTACACT TOTATAAGTA CCGTTTACTT CATGGCATGA ATAAATGGAT CTGTGAGATG AMBETTEMBE OCCUBETEET GETEMTEMET AMECAGATTT ACTIOGACTA CATOTGAAM GGCTGATGGG CTTCATCGTT TATAAAATCC GGGCTGCTGA TAAAAGAAGT AAGGCTTTGA NGAGGTOCGT TCTGAGCCGT CTGTCCTGCG CCAAGATGCT TCAAAGTATT ATTAAAAACA 420 360 300 240 180 120 60

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GGATAGAAGG GITTIGCAATG CCATATTAIT. GGTGGAGGC

ATGGCTTGCT GAATATCTTT ACCAACATCT TGAATATATA AGCAAAAAGA TAAAGCTTGG GTGGAATATC ATTITTAAAAT TITICITICACC TACTOTOCAA ATATIGIAAT GCAAAAGTO TIMATITICI GEICATIGIT ICICIICGAL AMAITIMIT

1440 1500 1560 1620 1680

GOTTTTGAAA TOTTTTTCAA ATATGTGAAA TOTGAAACTG ATTAAAGAAA ATGTATTIGTG ATTGAAATTA TTTTGACCTC

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ATTICITICCAG GGATTCTAAT ATTITATITAA G

ACTICTICAGTA GATAATTITOT CATGCAGGGC ATGCAATCAG AATCTCACTG AGCCACCAT	480	CTTTTAACTG AAAAGGGATG GGATAGAAGG GTTTGCAATG CCATA
CATTGTGAAA TAATTACCTC AGTTGTACAG GACTTGGTGA TCAGGATCCA GGCACTCACT	540	
TGTATTCTAC TGCTCAATAA AGGTTTATTA AACT	574	5 TTCTAGTGTC CACAAGATTT ACCAAAAGA TAAAGCTTGG GTOGA
		GITCARGITC IGITCIARAF TITCITCACC TACTCTCCAA ATAITK
(1) INFORMATION FOR SEQ ID NO: 271:		10 reagnasta titognagna tinatititica gescatigit icteri
(1) SEQUENCE CHARACTERISTICS:		TCATTAAATA CTTRTTAGAG GGTTTTGAAA TGTTTTTCAA ATAIG
(A) LENGTH: 1731 base pairs (B) TYPE: nucleic acid		CIGICITITA TAITAAAGIA AITAAAGAAA AIGIATIGIG AITGA
(C) STRANDERESS: double (D) TOPOLOGY: linear		O CACAAGATGG CTCTATGAGT ATTCTTCCAG GGATTCTAAT ATTTAI
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 271:		
GCTGCAAGGT GCGCTTCGTG CCGCTGCAGA TCCAGCTCAC TACCCTGGGA AATCTTACAC	09	20 (2) INFORMATION FOR SEQ ID NO: 272:
CTTCAAGCAC TOTOTTTTTC TOCTOTAATA TOCAGGAAAG GTTCAGACCA GCCATCAAGT	120	(1) SEQUENCE CHARACTERISTICS:
ATTITIGOGGA TAITIATTAGC GTGGGACAGA GALTTOTTGCA AGGGGCCCGS ATTITIAGGAA	180	(A) LENGTH: 1310 base pairs (B) TYPE: nucleic acid
TTCCTGTTAT TOTAACAGAA CAATACCCTA AAGGTCTTGG GAGCACGGTT CAAGAAATTG	240	() () ()
ATTTAACAGG TGTAAAACTG GTACTTCCAA AGACCAAGTT TTCAATGGTA TTACCAGAAG	300	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 272:
THGLAGOGGC ATTHGCAGAG ATTCCCGGAG TCAGGAGTGT TGTATTATTT GGAGTAGAAA	360	 стесттямова мандального намотноско нанаменно слитет
CTCATGTGTG CATCCAACAA ACTGCCCTGG AGCTAGTTGG CCGAGGAGTC GAGGTTCACA	420	TCACTATGTA GTGGAGGGC AGACACCCTC CCGCAAATTC TGGAAG
TTGTTGCTGA TGCCACCTCA TCAAGAAGCA TGATGGACAG GATGTTTGCC CTCGAGGGTC	480	35 crassociam ascociagae rocrasticee esserticase teacte
TOGETCRARC CHOGGANICAT AGTGACCACG AGTGAAGGET GITCTGCTTC AGCTGGTAGC	540	CTGCCGTCGC CATOTTIGGC TGCTTGGTGG CGGGGAGGCT GGTGCA
TGATAAGGAC CATCCAAAAT TCAAGGAAAT TCAGAATCTA ATTAAGGCGA GTGCTCCAGA	009	AAGTGGCAGA GGATAAATTT GTTTTTGACT TACCTGATTA TGAAAG
GTCGGGTCTG CTTTCCAAAG TATAGGACAT TTGAAGAACT GGTATGCTAC TCACTGGTGA	7 099	40 тосттитат ссгоссааса атессатите стемесами езсмос
AGGACAGTCA GGTGAAGGAC TGTAAGGCCA CACAAGCTCT TCTTATCTCT ACTAGAATTA	720	CTIATCCTGA TTCAAATGGA ATGCCAGTAT GGAAACTCCT AGGATT
AANTOTTAAG TCAAAAAGG CTCCTTTTTT GOGCCTCCTA GTGAACTTAA CCAGCTAGAC	780	45. ACCAAGIC CATCITCAA AITICAGGIC TIAAAICIGG AGAAGG
CATTIGACIA CCACCATITA GITACAAACG TCAAAGGCIT CCGGIGCIGC TIACCITICCI	840	TIGGAGCCAT GAATATIGIC CGAACTCCAT CIGTIGCICA GATIGG
TITITICITIAA TOTOCITITTA ITTAITIAAA AAAATTACAA TGAAGATGCC TGITITIGICT	006-	TATTAGACAG TATGGCTCAG CAGACTCCTG TAGGTAATGC TGCTGT
CTACTOTOTA CTCTGATGGT ATCTTTCCAA AGTGCAGACT CTTGTGAAGT TTTCTTAAAT	096	50 catteactea stecacaea aagatstiss acaaitticta caaitt
TOTICACTIT AAAGAAATG ACGIACCAAC AATGATTING CITTIAIAIT ACTOIAAGAT	1020	CTOTOTOTOA GGCCCAGNTG ACACCAAGCC CATOTGAAAT GTTCAT
GITATAATOT TAATOTOGAT GIAGTECTTT TACTITACAG ATTGATTGGA ATAAGATTAT	1080	55 TICHGGAAR GERAIGAGG AINTHCHGIC TCCAATAITA AGGTI
TOCATATGAA TITTACCCACA GGACTCTGAA TCATOTTACC CACTCCCCTC ACAMTGTTGT	1140	TCTATTITICT CTATGAATAT ATTCCTTTTT TGACATTTAA ACAIMI
CCACTTAGTG AGTTGCATTG ATCTATCGGT ACCAAATGAT GTTGAATAAT TACATATCTT	1200	CATCAGCACT GCATGCCATT AAAGTATGTA CTATAGAGAT CTGATG
TCTKGACTAT ACTGATTICT TATTITIGGTC ACTAITACTA AAICTCTGTT AATAITCTCT	1260	

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AGAGTICGCG GGGCAGAGG CATICTIGCC GCTGGCCCAG

AGACACCCTC CCGCAAATTC TGGAAGGTTC TTAGTCTCGA TOCTAGTOGC COCCTTCAGG TCACTGCCGG CTGAACGGAG 240

300 360 420 480 540 900 999

GITTITICACT TACCIGATTA TGAAAGTATC AACCATGITG ATCCCATTTC CTGAGGGAAT GGGAGGATCT GTCTACTTTT ATGCCAGTAT GGMAACTCCT AGGATTTGTC ACGAATGGGA ATTICAGGIC TRAATCIGG AGAAGGAAGC CAACAICCIT

TECTTEGREG CEGGGAGCT GGTGCAAACA GCTGCACAGC

180

840 900

720 780

ACACCAAGCC CATCTGAAAT GTTCATTCCG GCAAATGTGG

AINTICIGIC TCCAATAITA AGGCTITITTA TAACTGAATA ATTCCTTTT TGACATTTAA ACATATTCTT TTATTGTGAA AAAGTATGTA CTATAGAGAT CTGATGAGAA ACAGTTCTTA

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CGAACTCCAT CTGTTGCTCA GATTGGAATT TCAGTGGAAT

CAGACTCCTG TAGGTAATGC TGCTGTATCC TCAGTTGACT AAGATGTTGG ACAATTTCTA CAATTTTGCT TCATCATTTG

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SS S TACCATCTAT GAGAGTAGTT TATACTGCAC TGTGTACATG AATGGCTAAT GAATCTATTT GCCCTGATGT GGAGTACCTT GCCAGCATCT GCTGGGGTGA ACTITATTIT AGCCCTTCCC CATOCCTAGG AAGAGTTTOT TGAGAAGTOG TACCATOGTG TAGCATOGGA GAGCATTOGG OCCIATOTIT TAGGGACCAC TATTAAAGCT TATAAATATT TGTGTATTTT CATTTAGAAG CCCTAARTAT TITGITRIAT TOTCGCCRIT ATGRATITRI ARAGRCAGGA ARATRIRGIT GGWTATATGG CCTTGGCGAG TTCCCAGCAC AGCGCTCTG GAAGAGGCAT GAGGCATTTC TTTCAGGAAA TGRICATTAT CCCTOGAGAG GGGCTGCTGT GCCAGCTTOG GGAGGGTCTG GGATGGGGCT GCCCCTGATG 3 ATCTGCTCAT CTGANGAATG AGGAAATAGG AGTGAATTTG ATNITTCCTA GGTCCNTCTA AATGCACTAG GTTTGAAITT GGCATAATGG TAGCTATGTG ACCCTGAGCA AATTTCTCTC TOCAACTITO COOTSTITTA TAGATATITO TITTCACTIT GASTATOCTA GAGATGGGAG 2 GCGTAATTAT TACAGAGGCA GTCCATGTGC ATTGT TCAGCCAGAA GGCATTCATT AAGTAAGTCC TGACTTTGTG CCCAGCTCTG TTGTTOYTCT TATGAAGAAC AGAGGAGGG TGGGCAGGTC AGTGATGTCA GCAGTGAGTA TGACACCCAT AAGGAATTCA TGAAGAAAGT AGAAGAAAAG CGAGTGGACG TTAACTCAGC INFORMATION FOR SEQ ID NO: 273: INFORMATION FOR SEQ ID NO: 274: Ξ E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 273: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 274: TCCAAGAAGA CCCGGGGGGG GTGAAAACCC CACACTTIGA CITIGCTACC AIGGGCIGIG ICTANGNACG TATATATGCI ACTORGRAGG GGCARAGGRC GCTRGKTTKT AGWTARCRCG GRACCTCARA SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 515 base pairs (C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 2995 base pairs TOTOGACTAA TOCTCACOGO TOTTATAGGC 1320 1260 1200 1080 1020 240 515 480 420 360 8 180 120 8

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6 35 30 25 20 2 8 S 8 2 0 S CATTITIAGEA TECAAGITICE CEACEACEAA GITAGAGATG ACTOCTOTOG CTGACATTITI GATGACTOGC ANACAGOOTG ACGTGGACOG GGTCACCANG ACATACANAA GGANAMACAT CCAGCAGCGT CTTGAAACGG CCTTGTCAGA ACTGGTGGCT AATGCTGAGC TCCTGGAAGA CAAGGATGCG TATCGACCAA CAACCGATGC AGATAAAATC GAAGATGAGG TTACAAGACA ACTGAATGAT GCCTTGGATC GGCTGGAGGA GTTGAAAGAA TTTGCCAACT TTGACTTTGA CAACCAGCIT TCTGCCCGCT GGCAGCAGGT GTGGCTGTTA GCACTGGAGC GGCAAAGGAA TCAGCCAACC CCTCCTCCCA TGCCAATCCT TTCACAGTCT GAAGCAAAAA ACCCACGGAT CCCGCAGAAC ATTGACCGAG TTAAAGCCCT TATCGCTGAG CATCAGACAT TTATGGAGGA ACACTIGGATIC ACCATICATICO GAGOTICGOTT CGAGGAGGTO CTGACATICGG CTAAGCAGCA AGTAGCCATG GGAGAAGTCA TCCTGGCTGT CTGCCACCCC GATTGCATCA CAACCATCAA CIOGIDATAC CAGCANTIAG TICTICCCCG GCCTCCACAG GIGCCAAAAC TAAICGGGCA CATCAGGTAG CAAGTTGAAA CGACCAACAC CAACTTTTTCA TTCTAGTCGG ACATCCCTTG GOTOTACATO CATOCOATOT TOTOCAGOCA COCCAGOCAG TOGAACCAAG GITTATOCOAT CATCHICCCG GGCAGCTICC CCTACTCGTT CCAGCTCCAG TGCTAGTCAG AGTAACCACA TACCAGAGGG AGCATCCCAG GGAATGACCC CCTTCCGCTC ACGGGGTCGA AGGTCCAAAC GCGCAACCUT GATOOTTCGC GTTGGTGGAG GATGGATGGC CTTGGATGAA TTTTTAOTGA CCOGINCINC CICOGCAAIC AOTINGOGA TICICAOCAG TIGCOGCIGG ICCGIATICI AGTGGCTCAG TOCAAATOTO CAAAAAGGTT TCAGGTGGAG CAGATCGGAG AGAATAAATA CGACCGAGAT GOGGATGOTT ACATTGATTA TTATGAATTT GTGGCTOCTC TTCATCCCAA CTTCCGGCGC ATTGATAAGG ACCAGGATGG GAAGATAACA CGTCAGGAGT TTATCGATGG TGTCTGGAGG AAAAAGTATA TGCGTTGGAT GAATCACAAA AAGTCTCGAG TGATGGATTT AGAGCCTACT CACOCOCCTT TCATAGAGAA ATCCCCCAGC GGAGGCAGGA AATCCCTAAG ACTICIOSCA ISGAICCAGI GOSCIGAGAC CACCCICAIT CASCOGGAIC AGGAGCCAAI CANACCTTCC ANATCCCAA CCATOTCTAA GAAGACCACC ACTOCCTCCC CCAGGACTCC OCCAGCAGOC GOOGAGGAAG TGACGOTTOT GACTITIGACO TOTTAGAGAC GCATTGOTTG GACCCTAAAA AGTCTGCCAG TCGCCCTGGG AGTCGGGCTG GGAGTCGAGC CGGGAGTCGA AAAATGATCC CTGCCGAGCA CGAGGTAGAA CTAACATTGA ACTTAGAGAG AAATTCATCC AGGICCCAAG CGATAACACI GICTAAGCAC CCCCAAGCCA CTAICCACIT IGAAICCIGC TICCGACACI TCAGAAAGCA GCGCTGCAGG GGGCCAAGGC AACTCCAGGA GAGGGCTAAA ICCATACATT GOGTGTATAT TTATTCTGAA COOGAGAAGT TATATTGTTA AAAGTGTAAA 1140 1080 1020 1500 1320 1260 1200 1860 1620 1560 1440 1380 1740 1680 960 900 780 720 660 600 540 480 420 360 240 300 180 120

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	09	300	CAGAAATAC CCACCAGTAA AOTITITATC AGAAAAGAT CGGAAAAGAA TITITGAMTAA
aaatattga		240	ITTGAAGAG ATGGTTGAAC CACTAAGAGA GAAAATCAGA GATTTAGAAA AAAGCTTTAC
OTTGGGGGGC ACAAAGTINA CAIATTCTTG GTTAACCATG GTTAAATATG CTATTTTAAT		180	GCAACITIC GITAATATGA GGTCTATCCA GGAAAATGGT GAACIAAAAA TIGAAAGCAA
CATGATICAAG GATATITTGAA ATCACTACTG TGTTTTTGCTG CGTATICTTGGG GCGGGGGCAG	55	120	COCAGAGG ATGAAGCTGC TGCTGGGCAT GGCCTTGCTG GCCTAGGTGG CCTCTGTTTG
ACACTITITIC CAAGGICIAC TITIGAGITIC CAAACTICAC TITIGAAATAT TOCTOTIGGT		09	GACCOGCO COSCICCOGO GANTGTOAG CAAGGCGCTO CTGCNINCOTO TCTGCCGTCA
AFTGATTTTA GOGCAGATAA AAGAATTCTG TGTGAGAGCT TTATGTTTCT CTTTTAATTC	50		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 275:
GTCCTTATAC TATCCATAAA GAAAAAYCCT AGCAGTATTG TCAGGTCGTG GTCCCCCCC			(D) YOPOLOGY: double
AACATTAAGC GGGACAAAAA ATGCCGATTT TATTTATAAA AGTGGGTACT TAATAAATGA	•		TYPE: nucleic acid
CATATETTET GACTETGETT GACAGITITA TITACISCIT TETTTOTGAA GCTGAAAAG	45		(i) SEQUENCE CHARACTERISTICS:
TOTAMANIAG COTAGATICCA TCTCTGGGTA TTTTCAMGTT TTTTTATCTT GCTGTGAGAG) INFORMATION FOR SEQ ID NO: 275:
TAMANGCICC CCTCANANAA CTGCAGATTT TGCCTTGCAC TTTTTGAATC TCTCTTTTTA	40		
TOTCHTGAG AACAAACTOG AATTTCATTC TGAAGCTTGC TTTAATGAAA TOGATGTGC		2995	nachara arabarara rararara rararactics rogogoccc octac
ACTIGANGGO ATGIATITITI GCCTTTITITI TGITGICGIT TAAAGAAAGA CITTAACAG	. 35	2940	agctetac caatgaactg titiagaaaca agacacactt tigtaitaaa attigetigea
AAAGGAAGGAA CTCGCCACAG CTGAACTCCT CACTTTTTAGG ACACAAGACT ACCATTGTAC		2880	CAGCICCG ACCATGITIGC TOTOTGATTA TCTCAATIGG TITITAATIGA GGCAGAAACT
GANTICERET ACCAGGAAA TAATCAGTAC ATCCCCAAAC CAAAGCCTGC CAGAATAAAG		2820	ICTGACCAA GTGACCATGA AAAGGGGCTG TCTGGGGCTC CTGTTTTTTA GCTGCTGTTC
mazzarzong mzernonem geargaagest tyaaacaaag caattoacta cytoograaa	30	2760	CAGCACAC TIGITAACCA GICCIGITIG CITICGICIT TITTIGIGGG TAATAAAGIC
GCCCAGGATG ACCACAGAA AAGAAAACCA GACATCAAAA AAGCAAAGCT GATGCTGGGG		2700	MGCCTOTT CTGAAANTST GGACGTAAGA CAAACACGTG CTCGTCCTTT AATGGAGTTC
TITISCICACI DARIDADADA CCITICITICAS AGCESANCIG ADAITICAGIT ICTCICCODA	25	2640	TITITIGCT ATTITITIAA ATGGTCGATC AACTICCCAC AAACTGAGGA ATGAAITICCA
AACACCAACG TCAGCACCCC GGTCAACCTG GGGAACCCAG AAGAACACAC AATCCTAGAA		2580	OCTITISCA AGGATAGGO TOSTIGGIGA CATIGUGAAT ITCAGAITIG ITTITATICAC
TCTCAGACAA GGGGGTTCCA GTACGTCAGC GATCTAGTGA ATGGCCTCGT GGCTCTCATG		. 2520	ATTIGITG CTCCCTATCC ACCTAGACA CCAGTAACTC TTGTGTTCAC CAGGACCCAG
GRENGCIACT TCATCCTGCA GGGGCTCCAG GGGGAGCCAC TCACGGTATA CGGATCCGGG	20	2460	ngaattea googtaaatg taagtgteca gaaaacgtea gaacatttog gottttaaac
CARTGOCCA GANTETICAA CACETITGGO CCACOCATGC ACATGAACGA TGGGCGAOTA		2400	ITTGCAATA CTGTCTTTGG ATATTGTTTC AGTACTGGGT GTTTAAAGGA CAAATAGCTG
GOCAAACOTG TTOCAGAGAC CATGTGCTAT GCCTACATGA AGCAGGAAGG CGTGGAAGTG	61	2340	kctaaagag agggaacctc atctaagtaa catttgcaca tggatacagc aaaaggagtt
CAAAGTGAGG ATTACTGGGG CCACGTGAAT CCAATAGGAC CTCGGGCCTG CTACGATGAA	ž	2280	TCAATTAG CAAGAACTGA GGGAGGGGT TTTTCCAATG TTTAATGTTT TGTGAITTTT
GOTGCCCOTC TOCTCCT80C CTCCACATCG GAGOTGTATG GAGATCCTGA AGTCCACCCT		2220	ntiaaaaa ataagaance ngcaatsitt aaggaachet itititishaa arcaggaga
ANGACATTAA AGACCAATAC GATTGGGACA TTAAACATGT TGGGGCTGGC AAAACGAGTC	10	2160	ntattect aaatgeadet tetttataaa ettgaettge tateteagea agataaatta
GACCAGATAT ACCATCTGGC ATCTCCAGCC TCCCCTCCAA ACTACATGTA TAATCCTATC		2100	NAGOCTAT TAATAGGGTT TCTGCGGGGT GCAGGGTTGT AAACCTGCTT TATCTTTTAG
GGACATGAGA ACTICCAGIT GATTAACCAC GACGIGIGGG AGCCCCTCTA CATCGAGGTT	·	2040	ngaaatat ttgagaaaa caagtgaaa ggtcagata caartgtgta ttaaaaaa
GAGGTGACCS TGGTGGACAA TTTCTTCACG GGCAGGAAGA GAAACGTGGA GCACTGGAATC	•	1980	igaatatti atgiagataa aattigocic ciggiaacce igtaatogat ggggoocrga
CAGGAGGOG AGKGTTOGTG GGCTCCCATC TRAACTGACA AACTCATGAT GGACGGCCAC		1920	BATAATIG IGITARGAAG CIGCCITATI ITITITICITI TIGIAAGITA CIAITITICA

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20 5 6 ઝ 30 25 5 S 45 SS 8 TITIAAAGTCC GITGCATTGA AAATAACAAA CAATATCAAT GITTIAATCA AGGATCTCTT CCCHGOTTGG GGCACACGAG GAAATCGTAG TCGGGGAAGA CTCTACTGAA TAAGACATCA CAGCAATTTG GGCAACTTTA ATTATGAGCA GAGAGGAGCC TTCAGGGGAA GTAGAGGTGG CCACATTCCT CCTTCTTATA AGAGCACAGT AACACTATCC TGGAAACCTG TACAAAAGGT AACTTOCCTT AGCTCTCCAG GGTNAAACGG GTGAGNCCTT AAAAACAGAA GAGAACAAGA 2 AGGAGAAAGT AAGTIGCITT GCACCGCCIA CITAATICIT TICCATATAT IGIGATACAA GTCATAAAAG CAAAATACTT ACATAGCTTT CTTAAAATAT AGGAATGACA TTACATTTTT TITTOCTICTT COCTOTTAGT TITTTACCCA ATATATOGAG AAGAGTAATG GICAATUTTA TCAAATTOTG AATCTTTIAA ACATCTTGAT AATTTGTTGT TGAGAGCTGT TCATTCTAAA ATATTTTTOG ANGCTTNGTC TGCAANCITG ACTIGNTTTT GCAGTAICAT TATTCAGACT TIGIATOTAT GACCIACTIT IGIAACAGAC CAIGGIIGIG ICCAAGGIAA AACCACAGIG GOGGATOTICE CTTEMAACAG ACTOCTOCCT TCAGCTMAAA ACTEMATOTE CTTEMACCT GCATTOTICA GCATIGICAT GAGCTIAATA TACTTAAATT CTACTACTCA TIGGATIGCO TICAGCAGGA CCAAAAAGAG AIGCCAGGCA GAITITATAAC CCICCCAGIG GGAAATATAG TGAGATTGGG CAAAAGAGAG CCAGTGAAGA TACAACTICA GGTTCACCAC CCAAGAAATC GACTACACAA CATAAATCAC TITITIAAATT CCAGGAACGG GTAGICIGAC ACGGIGATTA ACATITIGIT TIMATIGITI AMIAMAGCIG CIGGGCAGIG GIGCAGCAIT CCIACCIAGI ATGTAATGAA ATTCAGTCIA GTICTGCTGA TAAAGATCAT CAGTTITGAA AGGTTACTGA GGATTCTAAG TTAGITGCAC TTACATGATT ATTGTTATTT AAAACTAAGA ATAAAGGCTG ACTITIGAAT ATGGAATCIT ACTATITIGAA TAGAAATGIG TATGTATAAT ATACATACAT CATTITICADA GATADATIGG DATIGCIGIT GOIGDADATAD CANCCADARI ACIGARICIG TOOTTITIGAG GOTGAATOOG TTATTAACTT GITATITIAGG ACATAACCAT ATATOTOTOT GTGTGTGTGT ATATATATAT ATATGCATGC ATGTACATAC AGGTTTCTAC AGGAAGAGAT GGTATAATTT ACAATTTGGA GATTTAATAA INFORMATION FOR SEQ ID NO: 276: Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double (A) LENGTH: 2436 base pairs TOPOLOGY: linear TITITIACICC CAGTAGCAAG TGTGAAACTT 1440 1380 1320 1260 1200 1140 1080 1020 240 180 120 780 660 600 540 480 420 360 300 960 900 840 720 8

> 30 23 20 15 5 S GAACCTAAAC TCCCTTGGAA TCTGAACAAA GGAATATAAA ATTGCCATTT GAAAACTGAC TAATGICATG GGCATTTITA GTAGCATAGA TATAAGTTAG CCAATAGAAT TITTAGGTTA AAACAACAGA TGGGGGGTTT GTGGAGTGTT CCAGGGCTAC CCAGAAAAAG TGACTTGATA ACATGGTACC AATAAGTAAG GGATGCTCTC GGTCAGCTTG GCTATGGAGT GGTGGCAATA GCACATGTGT GICTICATIA CCATIACCIC TACACIGCAG TGACATTTCT ATTITIOTIT CACAGITAAT TOGGTTTGCT TITGCCACTT TCAAGATTTT AACTTCTCAG GTTATTAATC AAAATTATTG GATACIGAGI IGACIGITEC CITATECETE ACCETTECCE TICCETTECE TAAGGCAATA CTITATAAGC ATATTIGTAA ACICAGAACT GAGCAGAAGI GACTITACIT TCICAAGITT TITGAAATAA ATTICCTTIT GTAATIITAA AAAAAA TITITICITIT GCAAGACACC TOITTAICAT CITOTITIAAA TOTAAATOTC CCCTTATOCT GTGCACAACT TAGGTTATTT TTGCTTCCGA ATTTGAATGA AAAACTTAAT GCCATGGATT AATTATAGAG ACTACAGCTA AATAAATTTG AACATTAAAT ATAATTITAC CACTTITITGT GGACCTCAGA GATAGATICAG CCAGTGGCCC AAAGCCATTT CAAGTACAGA ATGGGGAAAA TAGTTCTGAA AGTAGICIGA TGTATTTTIC TGAGGAATAG TTTGTGATTC CAATGCAGGT CTICCCICCC CAAGITITOCT ATTCAAATCA ACTOCCIGAA AAGAAGCAAA ACTCCTTTAT TAGAATTACT CCCTTTGTTC TGCATTTGAA TGTTTCGTAT ATCTCTAAAC ATTCCAAAAG ACCATGAGCT AGGCTAGAAT GATACAAGTG AGCAAAAGTT . 1680 2160 1740 2436 2400 2340 2280 2220 2100 2040 1980 1920 1860 1800 1620 1560 1500

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 782 base pairs

(2) INFORMATION FOR SEQ ID NO: 277:

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

Ě SEQUENCE DESCRIPTION: SEQ ID NO:

277

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S 8 8 GCAGGGCARA GOTGGGAGTC AAACCCGGGT GACAGGTGGG TGGAGAGCCCC TGTTTGAGGT TCACTIONG CCTTTRATTA AATTCCTAAG GGGCCTGAAG AAGACATTTC TACTGCAGAG TCACAAGITT AGIAGICCCA AAAIGGGTIA TATCCCTICC CCCTTTACAT CAGAAICTIG GOTTAGAGGC ACTTGAGCAA GGCCCCCACA TCCCAACTCT GGGAGTTOTG OTGGGAGGAG GCCACTGACT TCTCCCACCC TTCTGTCTCC CCCATAATAG TTTATTTGGT TGGTCTGGAC GCACTICIGG GGGATAGGAC CAGACAAGAT AACAGGAGCT CACAIGGINAA GCAGAAGCIG NGAAANGGGA AAACAACAGA AGGAGGGGAT CAAAGATAGC TGATCTCACA TGCTTCCCAG 240 120 420 360 300 180 8

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WO	PCT)	PCT/US98/04493
	\$00	
	TOTGOTIANT CCCTCTCTGG TATTAGTTTT TCCCCTGGGA GCAGGAAGCC CTAGGAAGAG	480
v	вазактискав ветессенов взатеттисе тесстесеет веатваваса вавасалает	540
ר	ACCTICICIAC COCCTCCCTC AAGAATIGG CTTGCCCAGG AATGCCCACC ACACATACCC	009
	TUTTOTITIT THEMANCAA ACTUTIONIN ANTICOMOGO THEOCHOCON CONTOCHOCO	
10	CICICAACCI ITACITCIGA ITICIATITIC AIGGAAITIG GGAITGAAGI IAAACIAGA	720
	CAGTGCCGCC AACACCAAGT CTTGCAGGAA AAAAATACAA AGAAATTTAA CAAAAAAAA	780
15	AA	782 .
20	(2) INPORMATION FOR SEQ ID NO: 278:	
23 23	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 961 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: 11near	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 278:	
Ş	анаттесвае товнанств тостетваес свосвестте несытваест своснанает	09
2	GGACTACACC ATCGAGATCC CGGATCAGCC CTGCTGGAGC CAGAAGAACA GCCCCAGGCC	120
	AGSTGGGAAG GAGGCAGAAA CTCGGCAGCC TGTGGTGATT CTYTTGGGCT GGGGTGGCTG	180
35	CAAGGACAAG AACCTTGCCA AGTACAGTGC CATCTACCAC AAAAGGGGCT GCATCGTAAT	240
	CCGATACACA GCCCCGTGGC ACATGGTCTT CTTCTCCGAG TCACTGGGTA TCCCTTCACT	300
9	TOOTSTITTS GOCCAGAAGC TGCTCGAACT GCTCTTTGAT TATGAGATTG AGAAGGAGC	360
}	CCTGCTCTTC CATOTCTTCA GCAACGOTGG CSTCATGCTG TACCGCTACG TGCTGGAGCT	420
	CETIGLAGACE COTOSCITICT OCCOCTIGOS TOTOSTOGOC ACCATETITIG ACAGOGOTICE	480
45	TIGOTIGACAGE AACETIGOTAG GGGCTTCTGC GCCTTGGCA GCCATCCTGG AGCGCCGGGC	540
	COCCATACTIC COCCIPATIVES TOCTIONTICAS CITTIFFICATION TOTTICACOT	009
ç	CCTGCTTGCT CCCATCACAG CCCTCTTCCA CACCCACTTC TATGACAGGC TACAGGACGC	099
3	GOCCTCTCGC TOGGCCGAGC TCTACCTCTA YTCGAGGGCT GACGAAGTAG TCCTGGCCAG	720
	AGACATAGAA CGCATGGTGG AGGCACGCCT GGCACGCCGG GTCCTGGCGC GTTCTGTGGA	780
22	ITTOGRETICA TETGEACACO TCAGCCACCT CEGTGACTAC CETACTTACT ACACAAGCCT	840
	CTOTOTOGAC TICANGGGCA ACTGGGGCG CTGGTGAGGC CATTGGTCCA TCTCAMGTCT	006
	CONTRACTOR REPRESENTS ASSESSED CONTRACT DIFFIGURACE RESISTANCE	V 20

ACCCITAGAA GITITICAIGI GIAGIGIGCI CAAAAGACAA GGITACGGAG AAGGCITICCG

TOGITIATAT GCICAGACAA CAGGAAAGGG GAGTATATCT CICAAAGAAC IGAATGCCCG

35

TCTTGGGAAT AAGATGGACA GACCTGAAGC CATCAGTGAA GAGAGGTTGC GAGAGATGTT

GOCTTACTICA GAGATTIGAT TIGGTCAACAT GCATAACTIG AATTICAATAG ACTITITIGGTG

4

CTGGATGGCA CAGTACATTG ATTAACACAA ACTCACATTG GTTCCAGGTC TCAACGTTCA

GITATABAAC AGAIGITITIT TAGAITATITA ATAITABAIC AACITAATITI GAAIGAGAAT

TGAAAACTGA TTCAAGTAAG TTTGAGTATC ACAATGTTAG CTTTCTAATT CCATAAAAGT ACTTGGTTTT TACAGTTTAT AATCTGACAT CACCCCAGCG CCATTTGTAA AGAGCAACTT TCCAGGAGTA CATTTGAAGC ACTTTTTAAC AACATGAAAC TATAAAACCAT ATTTAAAAGC

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WO 98/39448

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961

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COCCETTIGO AGTICOGICT CCTGGTGTAC GOCCAACGCC AAGTAGGGGA TIGGGTTCCC TCCAGTCGCA GCCCTATCA ATTIGGATAT GTCCTTCATA TTTGATTGGA TTTACAGTGG

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 279:

15

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1228 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 279:

TTTCAGCAGT GTGCTACAGT TTTTAGGATT ATATAAGAAA ACTGGTAAAC TGGTATTTCT TGGATTGGAT AATGCAGGAA AAACAACATT GCTACACATG CTAAAAGATG ACAGACTTGG

2

TACAACTITIT GATCITGGTG GACATGITCA AGCITCGAAGA GIGITGGAAAA ACTACCTITCC

ACAACATOTIC CCAACATTAC ATCCCACTTC CGAAGAACTS ACCATTGCTG GCATGACGT

25

TOCTATCAAT GCCATTGTAT TICTGGTGGA TIGTGCAGAC CACGAAAGGC TGTTAGAGTC

AAAAGAAGAA CITGAITCAC TAATGACAGA TGAAACCAIT GCIAAITGIGC CIAIACIGAI

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240 300 360 420 480 540 9 999 720 780 840 900 960 1020 0801 1140 1200

TCATCATGTT AAATTTTTTA TGTACTTTTC TGGAACTAGT TTTTAAATTT TAGATTATAT

S

GICCACCIAT CKTAAGTGTA CAGTTAATAA TTAGCTTATT CAATGATTGC ATGATGCCTT ACAGTTTTCA ATAACTTTTT TTCTTATGCA AAGGTCATGC AATAAAACAA ACTCTAATGT

TTGGCAAAA AAAAAAAA AAANTCGA

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960

GCTCCAGAAA TAAATGCCTG ACAMCTCCCC ACAAAAAAAA AAAAAAAAA ACTCGAGGG

(2) INFORMATION FOR SEQ ID NO: 280:

3 SEQUENCE CHARACTERISTICS:

(B) TYPE: nucleic acid LENGTH: 1327 base pairs

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(C) STRANDEDNESS: double (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 280

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GOCGOGTIOCA GEAGATICGTG GGCGCCCTCC GCAAGGGCGS CGGAGACCCGG TTACAGGTGA TOTOGOGIOT COGGACAGGI GAGCACCOIG AIGAAGGCCA COGIOCIGAT GCGGCACOIG

6

TITICIGATIT TRACATGACC TIGAGCAGGT TIGCATATAA TOGAAAGCGA TOCCCTICIT COCTOCTICA COACTATTAC CCAATTGAGA TOGACCCACA COGGACOGTO AAGGAGAAGC CTTACAATAT TCTOGATAAT AGCAAGATCA TCAGTGAGGA GTGTCGGAAA GAGCTCACAG

300 240 180 120

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TAATCCTTGC

ATGTTGGGTT

360

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AGAAOTTICA GATAGECEAG GIGGTIAGAG AGICEAATGE AATGETEAGG GAGGGATATA TACCTCATAT GGTGGAATGG TGGACCAAAG CGCACAATCT CCTATGTCAG CAGAAGATTC

25 TIGGIGATAT CCIGGAAGAA ATTAICCGAC AGAIGAAAGI GIICCACCCC AACAICCACA 540

30 TCOTOTOTAA CTACATOGAT TTTAATGAAG ATOGTTTTCT CCAGGGATTT AAGGGCCAGC 660 600

ઝ TIGAGGGCAA AACCAAIGIC AICCIGCIGG GAGACICIAI CGGGGACCIC ACCAIGGCG 720 780

AGCGGCGGGA NCGCTAACAT GGACTCCTAT GACATCGTGC TGGAGAAGGA CGAGACTCTG ATGGGGTTCC TGGTGTGCAG AACATTCTCA AAATTGGCTT CCTGAATGAC AAGGTGGAGG 840

AGGCCCCTGA AGGCGCAGGC TCCNAAGKCC SCTGCAGGCC GTGGTGAGGA GGGGCGCCTC GATGTGGTCA ACGGGCTACT GCAGCACATC CTGTGCCNAG GGGGTCCAGC TGGAGATGCA 960 90

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CCCAGAGTCT GUTCCCCCGT GAACACAGAG CAGAGCCAGG GTGGCCAGCA GTGGCTGGGT ATCACCAGAG GCTTGAAGGA

CCTTICCGCGC CCCTCCGTCC TOCTTTOCCT GAGCACCTTC

5

ACCCCGCCAT GTGGCAGGGC GGATTGTCTA CTCCAGGGAT TITICTTCAAA ATTITTAAAC ATGGGAAGTT ACAGGCACTG TICCIGGIGA ACCITIGGACC ACAGCATGIC

CARACAAATA TAATOTOTGA AACAGATCAA AATTTTTAAA ATGAAAAAAA AGCTGCTCTG

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INFORMATION FOR SEQ ID NO: 281:

ATTICAGGGGA TGTGGGTCGG

GGTAGAACCT GGACCTCTTG GCCTGGGGGC ACATGGGATG

1320 1260 1200 1140 1080 1020

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X

SEQUENCE DESCRIPTION: SEQ ID NO: 282:

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(B) TYPE: nucleic acid (A) LENGTH: 2196 base pairs STRANDEDNESS: double TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 282:

1327

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CIGGCIGGCA TOGGICTIGC TITICCITTAT ATGACTIGICC TGGGCTITIGA CIGCATCACC

180

120

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OCTGAGOCCE TOOGTACCTE COGAGATGGA TOOGTOTOCT ACTAGAACCA GOOTGTGTT AAAGACTOTA ACATOCATGA GOTTGAACAT GAGCAAGAGO CTACTTGTGC CKSCCAGATG

TCAGCTATAA CTGGAATAAT GOGAACTGTA GCTTTTACTT GGCTACGTCG AAAATGTGGT

300 240

ACAGGOTACG CCTACACTCA OOGACTGAGT GOTTCCATCC TCAGTATTTT GATGGGAGCI

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CHAMAMACT CGAGGGGG TITITOTITIC CITTATGAAG ઝ

TIGIOGATIT TAIGIGICIG IGGCTIAATA AICATAGIAA CAACAATAAT ACCITTITICI

CCATTITICT TOCAGGAAAC ATACCTTAAG TITTITITIGT

TITGITITIG

ITTITTTIGIT

AAAAATAAA ATAGTCACAT TTTTAATACY AAAAAATGGA

799 780 720 660 600

TGATACACA ATACAACAAG AACAGCTCTG TGTGTGAGAA CTGTGGTTAC TTCCAGCAAC

AGACCITCIT CAACACACIC TACCATAACA ACATICCCCT TITCATCTIT TCTGCGGGCA

480 420

25

AGAGAAGGAT TOOTGGATOT AGOTGGTCAO GAOGATGTTT TOACCAAGGT CACAGGAGCA ACATTIGAAA CAGICIGCAC CITIGATACG GIATIGCATI TCCAAAGCCA CCAAICCATT tigeoteset gatosostig aagtitostt toottettot tieageeeaa tatstagaga

480 420

360

GCCCTTCAAC CCCACCTTGG ACTTGAGGAC CTACCTGATG GGACGTTTCC ACGTGTCTCT

TAMAAATCTA TICAGAAATT GOTCCAATAA TOCACGTGCT TIGCCCTGGG TACAGCCAGA

ATTICCAOCCA AAGACAITTIC AAGTIGCCTGT AACTIGATITTG TACATATITTA

300 240 180

COCTITIONIC GARGESTAAT TECHEGOSTE CITOTITITT GAGAGAGACT GAGAGAACCA TOGINTICAC ACCIDENTS ACCETENESS CENSECCIOS GETGITECCE GIENCIGIOI CIGCIGAACC CAGCCIGGGC CIGGATGCIC TGTGAATACA TTATCITIGGG 120

TUACCOTOCO TACAGOTOG AGOTOAGATG ACTOCOCOCT COACOGTOAC TOTOAGOAGG

60

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SEQUENCE DESCRIPTION: SEQ ID NO: 281:

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(A) LEXCTH: 799 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS:

508

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9448	

509 TITOSTITOGAA CAGGICTGAT CTCAGGATTG GCACAGCTTT CCTGTTTGAT CTTGTGTGTG ANGTHERSTAT TEALCHEAT CTCAGGATTG CACAGGCTTT CCTGTTTTGAT CTTGTGTGTG	360
ATCTCTGTAT TCATGCCTGG AGGCCCCTG GACTTGTCCG TTTCTCCTTT TGAGGTATC CGATCAAGGT TCATTCAAGG AGAGTCAATT ACACCTACCA AGATACCTGA AGTTACAACT	420 480
GAAATATACA TOTCTAATGG GTCTAATATCT GCTAATATTG TCCCGGAGAC AAGTCCTGAA	540
TCTGTGCCCA TÀATCTCTGT CAGTCTGCTG TTTGCAGGCG TCATTGCTGC TAGAATCGGT	009
CITTOGTICCT TIGATITIAAC 10TGACACAG TTOCTOCAAG AAAATOTAAT TGAATCTGAA AGAGGCATTA TAAATGGTGT ACAGAACTCC ATGAACTATC TTCTTGATCT TCTGCATTTC	660 720
ATCATGGTCA TOCTGGCTCC ANATOCTGAN GCTTTTGGCT TGCTCGTNIT GAITTTCAGTC	780
TECTITIONS CARIGOSCEA CATTAINT TECCGATING CECAAAATAC TETGOSAAAE	840
AAGCTCTTTG CTTGCGGTCC TGATGCAAAA GAAGTTAGGA AGGAAAATCA AGCAAATACA	006
TCTGTTGTTT GAGACACTTT AACTGTTGCT ATCCTGTTAC TAGATTATAF AGACACATG	096
TOCTTAITTT GIACTOCAGA AITCCAAIAA AIGOCTOGGI GITTTIGCICT GITTTIACCA	1020
CAGCTGTGCC TTGAGAACTA AAAGCTGTTT AGGAAACCTA AGTCAGCAGA AATTAACTGA	1080
THARTITICC THRIGITICAG GCATGGAAAA AAAAITIGGAA AAGAAAAACT CAGITITAAAT	1140
ACGGAGACTA TAATGATAAC ACTGAATTCC CCTATTTCTC ATGAGTAGAT ACAATCTTAC	1200
GTAAAAGAGT GOTTAGTCAC GTGAATTCAG TTATCATTTG ACAGATTCTT ATCTGTACTA	1260
GAATTCAGAT AIGTCAGTTT TCTGCAAAAC TCACTTGT TCAAGACTAG CTAATTTATT	1320
TTTTTGCATC TTAGTTATTT TTAAAAGAA ATTCTTCAAG TATGAAGACT AAATTTTGAT	1380
AACTAAIRIT AICCTIRITG AICCTRITGA TCTTAAGGIA TITACAIGIA TGIGGAAAA	1440
CANAACACTT AACTAGAATT CTCTAATAAG GTTTATGGTT TAGCTTAAAG AGCACCTTTG	1500
TATTITIATT ATCAGATGGG GCAACATATT GTATGAAGCA TATGTAGCAC TTCACAGCAT	1560
GOTTATICATO TAAGCTGCAG GTAGAAGCAA AGCTGTAAAG TAGATTTATC ACACAATGAC	1620
TOCATACAGA CTTCAAATAT GTCAATAGTT TGGTCATAGA ACCTAGAAGC CAAAAGCCAC	1680
ACAGAAGGGC AAGAATCCCA AFTIVACTCA TOTTATCATC AFTAGTGATC TOTGTTGTAG	1740
AACATGAGGG TGTAAGCCTT CAGCCTGGCA AGTTACATGT AGAAAGCCCA CACTTGTGAA	1800
GOTTTIGITT TACADATCAC TIGATTIAAC ACACICAGGT AGANTATTIT TATTITTACT	1860
GITTIATACC CAGAAGITAT ITCIACATIG ITCIACAGCA AGAATATICA TAAAAGIAIC	1920
CCTTTCAART GCCTTTGAGA AGARTAGAAG AAAAAAGTT TGTRIRITT TTAAAAAATT	1980
CITITIADANG TCACITITICA ACAICITICI ACCANGAITGG TACITITICCT TAACCGITIA	2040
TATICACTITI CATGARACT GCAATAGOTT GCTATGAGCA CITTCTTAT CCTTGGAGTT	2100

TARICCITIG CITCATCITI CIACAGIAIG ACAINAIGAI TIGCIAIGIT GIAAAAAICIT 20 TGTAAAAAT TTCTATATAA AATATTTGAA ACTTAA

2160 2196

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(2) INFORMATION FOR SEQ ID NO: 283:

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(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1185 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 283:

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120 180 240 300 360 480 240 909 099 720 780 840

420

TAAAGTOCCT TOOGTATAAA AATCTAAATG TCTGCGGTGT GATCAGTCAG GAGCACGTAA STUCCAGGET AAAGGAAAAG CITCACAAGG GIAAGAGCCA CAGAACCCIC GGCAAGAAAG GCCGGTCAGG GAGAATGAAT GGTACAGAGA GGAAAGGAAG GAAAGGGGGT GGAACAGAG TAGAAGGCAA GGAAGGGATG CCGCACTGGA GACCGATGGG GACACTCTAA TTGTGCAAGA GOGAGGANCT TECTTETAS ATSCTGAACA CAGCTAGTET GAACETTEET TGGAAAGTEE AGCIGITIGC CCATGCATAG GGCCAACTCT CCCTGCAAAG CAGCAAATGT GGCTTCTATC AGGAAGGAAA AGTATCCATC AGTGTGACAA GAGGTCACCT TCGAACTTGC ATGAACTCCT TOCOCACCA CAAAGAGTOC TOOTAGAAGT GAGGATOGCO TAGTOTTAGG GCTGTCGGTT TATAGAAGTA GCAGTACAAC ACTGCTGCTA GTCTCTGGAA TACAAACAGC ATTTGAAGTC CATCHOTICCA TATGAAGCTG TIGGAGTTTT TCCAGCGTAA GITCATGACC CAGACATGAA GGGAGATGCT GAGGGCAAAG TACACAGCTG TCAGGATGAT GGTCCCTTTG AACTTATGGA ATAGGAGOTT GACCAGGCCA GCCTGGAAGA CGAAGGTGTT GAAGAACATG AGGAAAATGA TGATGATETT GAAGAGGACT GCAATATCCT GGATGCACTG AGGGAGAGGY TTCTAGTTCC CHAICACHET TEGENICETT INGICACTEG GAGAICETTT GGGGGCTGGG AGGICETTET GCAGITIAAGG CITICIGATAA GGAAAGAGAG TCTGAACAGA GCACACACAT CTGGAGCTCC CTTTGTACT ATAANGGATE TEAACAAAGA YETGIBATITIE ATETIGIGGET CEATETICEE TENGGGICAA AGACTCCCCC ATTTATGGGA ACAAGAATTC AATTTATTCT CTATTTATAA AACATTTTT GIAGAIGITA AGCIGGACCT IGGCACGCCT CITIAACAIGA AGAGAICTAG CIAGACAGA AGGACTOGGG GATGCAGCAT CAGATTCCCAT CTTGAATTTC TGCTAAAATA ឧ 3 6 S 35 45 55 25

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1140 1185

TITIGAATGAG AGCTGTTTCC CTTGCTCTAA GGCAAGCACC TCCAA

1020 1080

(2) INFORMATION FOR SEQ ID NO: 284:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1634 base pairs

KITICACCGGG GGTC

CIGGOCOCGO IOGCICAÇAC CIGIAAICCC AGCAYITIOG GGAGGCCSAG GOCOGGGCGG IGAGITICCIO TGIGICCAAA ACTGAGGCAC CATGITICITT GAAAACATGC CACCTCAAGG

1620 1560

1634

6 ઝ 30 25 20 15 45 5 55 50 S CCACCACAAA TTOTOTACAT AGTCTTCAGA TGATACCACC AAGCTTGGTT TGTACAAAAG CAAGGTGGGA GTCTATTTTT GTACATGAGA TACATCACAC TARACARACA TITATIGUAGO CARCACTOOT TIGUAGATOOR GARACTIGAGO CACARTAGGO CCTTGCTTTC TCTTGGAAGA GGAAAGGACT CTGGTCAGGC CCAGGCTGAG TGAGATGAGC CICGITICCA AIGATAGAIC ACICCIGITG ACCIGGIAIG ICIGCITIGCI IGCIGCITIT TOGTACTO CAAAGCTGAT GTTCAGCTGA ATACAGATAC AAGAGCTGGT TCTAGGCCTG TGTTATATGT CATATTTAGC TTACCTGTGG AGCAGCCTGG TGCAGTGCCC TGTCATCAAG ACAAACCCAC GGTCCTNCTG ACCCAAAGCG GAGGGTAGCG GGAGGGTAGC AGGTGAGTTC CTAGGGCTGG AAATGATCTT 'AATTGTTATT GCCCACCCCT GGCTTTTCCG GGTAGAAAAT TCACAGTAGG TTATGACTIC CTCAAGAATA TGTAGCIGCT AGGGGGTAAA TCAAGGCATC ACAATTICTG TOCAGOTOGO TOATGGOOTT OTTAGAGCAG AGAGAGGAGT ATGTCATTIT ACTAAGITOO TOGITIOTOG CATTIGGGGA TAAGGIGCTG AAGCCAGAGC ATTIGCAGIT TGITTGAGGC IGIGITICCIC TGTTTGCCAC ATTICICITIC TIGACTIGIT CCICIGAAAG IGCAAGAGGC GIACACCIII CCCAAAIGIA GACIAGAAIC TICAGCOGGC AGGAATAGGC IGIGAATIGC TAGCACIFIT TITITITAAG CAATTACTIT ACATATOTOC AACCAGAGCA GCCACCAAGC ATTACTTAGC AGCAGGAAAA TGATTGTATT AATGATTOTT AAGAGAGAGT GCTTGGAACC ATGGGTTAAC AGGAAAGGCT ACCTAACTTC IOCAGGATOC CACOCACTOT ATAGITICIGO TITICOCAGAG AGGAAGAACT TITIAGAAACO (xi) SEQUENCE DESCRIPTION: SEQ ID NO: TCTOTOCCTO TOTOTOTOTO TGATAGICAC TCTTGCATGG CTTCCATGTC AAGCTGCACT TIGGGGCCAT CICIGCAGIA TIAGCCCCCT TITTGCTIGG AGIGITIGOT GITICITIGO CAGCIAAGIG AGGGICTIGG GAIGACTIGO AAGTATTAGT GAGTCTTCCT TATTAATATT TICATTICAG AAGACTGAAG GCATATATGT THIGHAITH OCCACTATTO TGAAGTGAGT CTGAGTTGTT TACACTGATG CCTTCCCTGC AGCAGATIGGA GTCCTGCAGC CCAGGAGACA CCCTGCATCC CTGCTAATAG (D) TOPOLOGY: linear (C) STRANDEDNESS: double (B) TYPE: nucleic acid GIGIGGGIAT GCATATAICT CICATCIGIA GITTCCAAGA AGCACAGIGT ATGCACCITC ATTTAAATAC ATCTGTGTGC STITTIATAT ATGACCTITG CCTTTCCCCA GCTCCCAACC OGTIGOCITACO 1140 1080 1200 1020 1500 1440 1380 1320 1260 660 600 540 480 420 360 960 900 840 780 720 300 240 180 120 6

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1260 1200 1140 1080 1020 960

900 840 780 720

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear TCACTGCAGT GTCAGGCCTT TAGATGGGAC CTGGGAGAGG CCCTGCGATT TCCCACTCCT GGGCATGGTG GCCCATGTAG cicicicico TICCICCICC TICITAGICT CCAGGAGATC Terecrose CCAGCGAAAA CATCCACCTA CTCACCATCC TITCICICIG

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420 360 300 240 180 120

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660 909 540 WO 98/39448

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%	WO 98/39448	US98/04493	:	PCT/US98/04493
	513		514	
	AATHGACCAC ATTGAAGGAG CACAATGCCC TCCTGTGTTG ATGCCACTTC CCAGGGTGGA	. 0211	TIMAMACTI ATMAMCTA	858
v	CACACTOGAA AAGAACCGAG GACAGGAAAG GATTGGGTAG GTGAAGGGGT CAGGGGACTG	1380	-	
n	GTAGTCACCC AATCTTOGAG AGGTGCAAAA AGCACTGGGG GCTACCCGTT AGCTGCATCT	1440	(2) INFORMATION FOR SEQ ID NO: 287:	
	GCCCTGGCTG TITICCCCGTT CATGTCACAA ACTCCCACTA CTATGTACCT GCAGTGGGGT	1500	(1) SEQUENCE CHARACTERISTICS: (A) LENTH: 915 base pairs	
0	TECHGAGATG GGGGAGACTC ANGTETTACT COCCAGGAGC TOCCAGGGGCC CAAGGAGGAG	1560 10	8 9	
	AATGCTGCCT CCTTTCAGTC TGGTCTACAC CCACTTTCTG GTAGCCTCTC TGCTTCCTGT	1620	(D) TOPOLOGY: linear	-
ž	AATTOTGBOT GITTITICCAG ACTOAGCTCA AATAGTGCCC CTCCTTAAGC CCATCCCTCG	1680	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 287:	
3	CCCCCAGCCT GAGGTGATCT TITCCCTCCTC TGAACTATTA GAGCAGTTAC TGTCTGTTCA	1740	GAATTCBGCA CGAGCGCGC CATGGCGCTC CTGCTTTCGG TGCTGCGTOT ACTGCTGGGC	09
	OTTOOTITIOG CAGGCACA CAGTGGCATA ANTICTATIG TITTICAACTC TGATT	1795	GOCTICTIVOS COCTOGREGO OTICOCCAAG CICTOGGAGG AGAICTOGGC TOCAGITIÑO	3 120
70		20	GAGCCGANTGA ATGCCCTGTT CGTGCAGTTT GCTGAGGTGT TCCCGCTGAA GGTATTTVGC	180
			TACCAGCCAG ATCCCCTGAA CTACCAAATA GCTGTGGGCT TTCTGGAACT GCTGGCTGGG	240
_	(2) INFORMATION FOR SEQ ID NO: 286:		TTGCTGCTGG TCATGGGCCC ACCGATGCTG CAAGAGATCA GTAACTTGTT CTTGATTCTG	300
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 858 base pairs	25	CICATGATGO GOCTATOTT CACCITGGCA GCTOTGAAG AGTCACTAAG CACCTGTATO	360
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		CCAGCCATTG TCTGCCTGGG GTTCCTGCTG CTGCTGAATG TCGGCCAGCT CTTAGCCCAG	420
30	(D) TOPOLOGY: linear	30	ACTIVAGAAGG TGGTCAGACC CACTAGGAAG AAGACTCTAA GTACATTCAA GGAATCCTGG	. 084
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 286:		ANGTAGAGCA TETETGTETE TITATGCEAT GEAGETGTEA CAGEAGGAAC ATGGTAGAAC	540
	TCTGCTTTCG GTGCTGCGTG TACTGCTGGG GGGCTTCTTC GCGCTCGTGG GGTTGGCCAA	09		009
35	GCTCTCGGAG GAGATCTCGG CTCCAGTTTC GGAGCGGATG AATGCCCTGT TCGTGCAGTT	120 35		099
	TOCTGAGGTG TECCEGATEA AGGIATTING CTACAGOCA GATOCCCTGA ACTACAAAT	180	ANGTIANGCA TAITAACAIT COTCANGICA TAIGAMAATA CAAAATAAGC AGAAAAGAAA	720
9	ACCIDIDADO TITICIDADAC TOCIDOCIDO GITIGODORIO GICATUDADOS CACOGATIGOS	240	TITAAAICAA CCAAAAIICI	
	GCAAGAGATC AGTAACTTOT TCTTGATTCT GCTCATGATG GGGGCTATCT TCACCTTGGC	300	TACCICTGAA CTITITICTG TGCCTTFAAA CAGATATATA TTITITITHA ATGAAAATAA	840
	AGCICTGAAA GAGTCACTAA GCACCTGTAT CCCAGCCATT GTCTGCCTGG GGTTCCTGCT	360	AACCADADA CCDATTITAT TICCICCITT TAAAACCITA TAAAACTATAA HAAAAAAAAAA	006
45	GCTGCTGAAT GTCGCCCAGC TCTTAGCCCA GACTAAGAAG GTGGTCAGAC CCACTAGGAA	420 45	-	-
	GAAGACTCTA AGTACATTCA AGGAATCCTG GAAGTAGAGC ATCTCTGTCT CTTTATGCCA	480		
ç	TOCAGCTOTC ACAGCAGGAA CATGGTAGAA CACAGAGTCT ATCATCTTGT TACCAGTATA	540		
3	ATATICCAGGG TCAGCCAGTG TTGAAAGAGA CATTTTGTCT ACCTGGCACT GCTTTCTCTT	009	(2) INFORMATION FOR SEQ ID NO: 288:	
	TITAGCTITA CTACTCTITI GTGAGGAGTA CATGITATGC ATAITAACAT TOCTCATGTC	099	(i) SEQUENCE CHARACTERISTICS: (a) LENTH, 1517 base pairs	
55	ATATGAAAAT ACAAAATAAG CAGAAAAGAA ATTTAAATCA ACCAAAATTC TGATGCCCCA	720 55	(A C	
	ANTANCACT TITAATGCCT TGGTGTAAGT ATACCTCTGA ACTITITITCT GTGCCTTTAA	780	(D) TOPOLOGY: linear	
9	acagatatat attititit aatgaaata aaacatata tectatitia titectecti	09 098	(*i) SEQUENCE DESCRIPTION: SEQ ID NO: 288:	

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6 ઝ ઝ 25 8 15 5 3 50 55 න TOGTTCCACT GGGGCAGCT AACGCTGAGT GACAAGGATG GGAAGCCACA GGTGCATTTT GOSCITGAGI CIGGCAAGGA ACCITGCIIT TAGCITCACC ACCAAGGAGA GAGGITGACA GCGTGGCTSC CCCCCAGGGC CACCGCTTCT TTCTTGATCC TCTTTCCTTA ACAGTGACTT GICCCCTAGG TATCAGCCIC ICTTACIGIA CICICCGGGA AIGITAACCI TICIATITIC ACTCAAGICT TCTCTAGTCA ATGAGGGGCA CCCAGTOCTT CTAGGGCAGG CTGGGTGGTG TGACCTTTCT CICICCICAT TICGGIGGAT GICCTTTCTG CAGCIGCCIT TCAGCACAGG TOTAGOGGG GAGCTICCTT GAGCAGIGGG CCCAGGCCIG GCCCICCACA CITCATICIC TAAAATGITA GAAGTATATA TATACATATA TATATTTCTT TAAATTTTTG AGTCTTIGAT ATCTACCCCA CCCCCTAGGA TCCAGGACTG GGTCAAAGCT GCATGAAACC AGGCCCTGGC AGCCTOTOCC ACCTOTOTAG GCAAGCTGGC TTCCCCATTG GCCCCTOTOG GTCCACAGOA CTGCCCACTG AGITTGGGGAA AGAGGATAAT CAGTGAGCAC TGTTCTGCTC AGAGCTCCTG AMPCANTATT CTTCCCTTGC CTGTGGGCAG TNGGAGAGTG CTCCTGGGTG TACGCTGCAG CICTOGCCCT CICCAGGGIG TITTICCACTA GICACIACIG ICTICICCIT GIAGCIAAIC TRACCTICCCC GCCCCCTCAC CAAGGCTGGG AACAGAGGG ATGTGGTGAG AGCCAGGTTC STOTETTOTE CATETTETAL CECECACECE CEATTACOOG TAAAGGRAAC CECAGACTAG GCACACHTIG GACCCAAGTA TOGGCCICTT CTGCCTAGTA CTGCCAAAGG GACTGTTAAG OTTICATITA AAGAIGITAA TIAAATGAIT GAAACITGGC TGTGGCTACT ATGTCTAAAA ATCCATTCCC TCTGCCCTGA AGCCTGAGTG AGACACATGA AGAAAACTGT GTAAATGAGC TTGACCTAGA GTAAATGGAG AGACCAAAAG CCTCTGATTT TTAATTTCCA GCACAGAGGA CAAAGGAGAA GGGAGGGTCT AGAAGAGGCA GCCCTTCTTT GTCCTCTGGG TCCAACATTA CIGGAACICI AICCIGITIAG GAICTICIGA GCTIGITICC CIGCIGGGIG AGCAACCCIG GGAANGGCIG GAGGNOGGAG AGAACCIGAC TICICTITICC CICICCCICC AAATTCCAGG AGCCCTGGG CAGGCCCTGG NCCCCAGTGC CAAGCCTCAG AGTAAGCAGA GTTGGGGGGA CAGGGCAGTG GTCTGGGCCC ACATTTAGAA GGGAAAATGT X SEQUENCE DESCRIPTION: SEQ ID NO: 289 AACAACACGC AGCTGATTTA GOGAGTGTCC CAGCCTAGCT GGATCAAGGG GAAGTACCCC ACCCTCTATG AAGACAGAAT CACTCTCTGC CATTCATTCT TTCAGACATT TTATTTGAAT TTATGACAGT GATGGGGATT AGCAGCIGCC TCACATIGIG TICICICCIG AGAITGGICCA GCICACATCC INICCIGCI TGACTGAGAT 1020 1200 1140 1080 1680 1620 1560 1500 1440 1380 1320 1260 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120

960 900 840 780 720

1320

1500 1440

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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E SEQUENCE CHARACTERISTICS: (A) LENGTH: 3865 base pairs (B) TYPE: nucleic acid

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2 25 20 5 35 30 3 6 S 50 CAAGGANGGG CGCTTGTTCA ANGAGCAGAA CTTCTTCCAG CGGGCCGCCA AGCCTCTGCA CCACCTCACT TGAAGCTTTG CCCACAGGAC AGTGCTGACA GACAAGAGTG GOCGACAGTG AMAGCAGOCO TOMGAMGACO AGGAMGAGCO CTOMGGTOMO CAGGGGTAGO COTOMGAMGA CCTTOTOGCA ACTAGTOGGT CCCCCGGGCT GCAGNAATTC GGGCAGTGGT TCTGNOTCTG GEAGGAGECY TEACGAGGGG GACCTTGAGT TYCATTAGEA TGGACCTGCA CAAGGGATGC ATGTGACAGC TGAAAATATC TITOTXGATC CAGAGGACCA GAGTCAGGTG CTCCACCCTC ACCTGTGACT CAGGACCACA GAAGCAAAAG TTCTCACTCA AACTGGATGC GAAGCTGAAG TCCTTCCAGA CCAGGGACAA CCAGGGCATT CTCTATGAAG CTGCACCCAC CCAGCTOTAG CCCTCAGAAG ACCAGGCAGA GCCCTCAGAC GCTGAAGCGG AGCCGAGTGA AAGATACTOT GAGTTOCTOT GAGAGATOCA AAGGCTOCGG GAGCAGACCC CCAACCCCCA CCAGTTAAAA TCTCCTCAAA ATGTTTGGAT ACCGCCCATT GGCCCCTCAC AGCCACGAGC GACAGCCTAC TIGACOGCCC CGCIGGCCCC CACATICCAC TGAACIGIGC GGAIGCCACA GATATETECCS TEATGCAGEC GECTECGGGG GACCACCTEC CTCCCTTTGA GTCAGCCACA GTCACCGTCA GCAGTTTOCA GTTTTCCACC TCCWCCCAGT TCCTCCGTGT GGTTGACCCA AGTIVOCCITIG GGATICATITI THATIGIHAGO TRGACITITOT CATGCCHGAA ACAAGGCING GOSTITICIOS CATOGACAAA TIGOCTITOCA AMAMIGAGGA CATOATGAAG CAAAAACAGA GOGCCCTCCC GCCGCRGYGA CCTCCAGAGC CTGGGYTANT GCATGCTGAA GTGGYTCTIAM OCTATOSCIT COCNTICCOC TATTOCCCAA GIGÓCAAACA COTOSCCIAC TOGOCTIGACG GOTGOTOGAAT GOOCTIGGAGT TOOTOCATGA GAATGAGTAT GTTCATGGAA TCAGTOGGCC CTGGATGTCA GCCCAAAGCA TGTGCTGTGC AGAGAGGTCT GTGCTGCAGG TTICGGIGIT CACCAGGACA AATACAGGIT CITGGIGITIA CCCAGCCIGG GGAGGAGCCI AGTEAACAAG TOGAAGAAGE TGTACTEGAE CECACTOCTG GECATECETA CETOCATGGG GTGACCCCCT CTCAGGCACA GCATGACCTC CTGAAGTCGA GCCTGCTTGC TTTGAACCTA ТТССАВАВАВ ВАВВАВА GGGAGGTCAC CTITIGGGIGT GIOLOGIGI GIGIGIGIGI GIGIGICIGI GIGIGIGCITI GGGACGGGIG GCGGTGTGCC CCCAGGACCT GTAAGTAATA AAATCTTTAT 1260 1200 1140 1080 1020

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	CAITIGGGAAA GIUGCCAACC ACTIGGIAGA CCACIAGGIT CICTGITITIC CCITCCCTIT	1740	CATAGTAACT GACTCGGAGT TCTAGAAGTT CCCATGGGGC TTTCATGTTA TCCAGGTATG	
v	CCTTTICAAA TCCCACAGIT TCCTGTIGGG GAGAAGCIGT AATTAGCCIA GTCCAGGIAC	1800	COTIGGAGGIC COCAAACAGA ATTOTTACCT CACACCCIGC CTTTAAGAAG TCTGCAATCT	
.	CAGATCCCAG CTAGGGGCGC ACCTGNCTTG GATAACTCCA AGAAAACCTG GGCACCAGTA	1860	THEACHTISGS CACAAAGTAA GCCACATGTIS GTTTGCCCGT GGTTGCCGTT CCCCAGTAAA	
	TITITICCAAT TATAAGSACT GTGGCATAAA TITITIAAATG AGTTATATTG AAACCAGATT	1920	TITITAAGITC COGCICCITIC AGIAICICCI TCAGCITICIC ITCCCCCAGA ACCICCTGCA	
Ö	TCTCCAGCTG CCAAGGGAAG AAGGTAGGGC TGGACTCCCT GCTGTGGCCC AGCCCTTGTT	1980 10	GOTTCOGGOT GATAAGONGC AGTITCTCTT CAGGGCTGGG AGGGTCCCCC ATGGTCCGCT	
	AGGGTTGCT CTCTCACTGC AGCCAGACAG GATGATCCTG GGTTCTGGGG AGGGTAAAGCT	2040	ACCICIACIT CCCCCGCTCA GCCCGGCACC AGAGCCCCTT CCTGGGTCAC CGTCGCCGGC	
v	GCCCCTTGCC CAGITYCTGCA CCCAATAAAG AGTCCAAACC CGCTGCTTCC GTGTCCTGAG	2100	GCGTGCCGGG AACTGTCACG CGAGT	
2	AGATOSOTIVA ATOSOTIGATO GATOGACAG ACTOMAGAGA CAGOAGATGA CTCADTOSTO	2160		
	GAAGAAGGG GGAAGATGCT GGCTGGCTA GCTAATGTTC CCCCCTTTCA GCGATTTACA	2220	:	
Ó	GGAAATGGAG CCCHGCTTGG TCATGAAGTT GGTTTGCTTC CACTGTGGGA TGCACTCCTC	2280 20	(2) INFORMATION FOR SEQ ID NO: 290:	
	AGANAITITIG ANGTCAGCCT GCAACTICTC GAAGACTITIC ITCTITGGGCT TGAGCTCCTC	2340	(i) SEQUENCE CHARACTERISCIS: (A) LEWOTH: 1910 base pairs.	
Ý	ATCHGGTIGG CCCTTTICAT AGCCCTTCAC AAACACGTGC TCACCAGGAG CAGAGCCTGC	2400		
3	CEGAGGGTCC AGAGGTTCAA CTGGCGGTTT ATCCCTTCTA TAGAAGCACA CAGAAGCATG	2460	(U) IOPULOAT: AIREAL	
	CETTOGGACT CARETECTET CATETYCTGG GOTTTCAGGT TOCACAGAC CACTACCAGC	2520	(X1) SEQUENCE DESCRIPTION: SEQ 1D NO: 230:	
0	CTOTOCTICCA CITOCTCCTT GOSCACGAAC TOTIACCAGGC COCTCACCAC AGTOCOTIGOT	30	AGGANGGA GANGGGG TCTOCCCCCG GCCGTTACCC AGANGCCACC GGACGGCLACC	
	TCAGCTTCCC CCACGTCAAT CTTCTCTACA TACAGGCTGT CTGCATCTGG GTGCTTCTCC	2640	ACSGAGTIGGG CTGTCCCCCGA GCCCAGCCCC GAGCGAGCCC CCCCCCGCC CCCGAAGAAC	
	ACAGIGATGA TITTCCCCAC ACGGATATCC ACCCGGATG GGATGACCTC CTCTGGTTCT		GOSCCTYCCA GCCAGOCCGA CTCCTAGGAG GAGGGGAGGC GGGAAAGCAG CTCAAGCCTC	
0	GAATICITIGG CAGGCCTITIG GCCAITIGGCT TCTGCTITIGA GGGAITCTGGG TAGGCAGCGC	35 2760	אבסכאסכסכ בושככסככאם כככספכבאבד כככאפסבדכ וכספאבדכה פכספדסכדכ	
	TOSCCAGTIT TITICAGGSCA GGGSTATIFA ACTITITOCOG GATIGGATOC AGCAACTIGF	2820	CTGGGAGTCT CGGAGGGGAC CGNCTCTGCA GACGCCATGG AGTTGGTGCT GGTCTTCCTC	
c	TEACHTOCAE THEADLACAA THETTERACTE PHYCALEGIAGE TO TABLADETHE	2880	TOCAGCCTICE TOGCCCCCAT GETCCTICGCC AGTICCAGCTG ANAGGAGAA GGANATICGAC	
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'n	GGAILGALAAA CILGGAACITA AGGGGAAAAAA GGALAIGCIT GAIGAAGGAAL AGAACLUCAT	3000	CCCCGGGCCC CAGAGATGA GGAGCCCAG GTGGAGAACC TCATCACCGC CAATGCAACA	
	TETTETCEAC ATTICCTOSC TCACAGAAGG CCTTCTTCAG TTTTTTCTTC ACATCCTCCT	3060	GROCCCCAGA ANGCAGAGAA CTGAAGTGCA GCCATCAGGT GGAAGCCTCT GGAACCTGAG	
	TOGGATCAAG GAGATCAATC TTGGACTCCT CTTCTGAAGA GCTCATTTTG CTGCCTGTTA		GCGGCTGCTT GAACCTTTGG ATGCAAATGT CGATGCTTAA GAAAACCGCC CACTTCAGCA	
ġ	ATCCTGGAAC CATAGGATTC ATCAGATGGA CCCGTTTTGA ATAGCCAAGT GCAGGGAGGT	3180 50	ACAGCCCTTT CCCCAGGAGA ACCCAAGAAC TYGTGTGTCC CCCACCCTAT CCCCTCTAAC	
	ACTICICISC AAAGGIGAAA ATCITICICT GATCAATGCC TCCAAATTGG GCALCTACTT	3240	The second section of the second section of the second section	
v	TIMMINCIC TICHICCAM GOCTGCAGTC CGGGGTATAM GAGGCCACTC AGCAAAGGGT	3300	ACCATTUCTO CACCIONATON TOCANCINAL AUTHOUNTE CUACHOCAGO, CINCOGNICO	
,	GCTCCACCTG CTTTACCACC TCAGCTCCAG CCTTCTTGGA ATCGTGCTGT GTGACCACGG	3360	SCCALCICC CSIGNIGISI GIGISISI GISTOLOGISI GACISIOISI GITTOCTANI	
	AGGAGAGTET GTACACATET AGTGTGTACT CTTTGCTGAG CTGGTAATCA GTGCCTTTGA	3420	TORGOTOTT GIGGOTACTT GITTGIGGAT GGTATTGIGT ITGITAGIGA ACTGIGGACT	
.0	TSAACTIGAG CITCTCCAAG GGCACACCAA TGCTCTCCAG CATTGCTTTG ATCACATTCT	3480 60	CACTITICEA GGAGGGGT GACCACATG GCCATCTGCT CCTCCCTGCC CCGTGGCCC	

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GOTTTGCAGC ACTITISTICAT CATTCTTICAT GGACTCCTTT CACTCCTTTA ACAAAAACCT

1200 1140 1080 1020

TTAGGGATGC GTAGGGTAAG AGCACGGGCA GTGGTCTTCA GTCGTCTTGG GACCTGGGAA

TOCATOACOT TOTOCTOCTA GUAGGOTGOT TOTTOCCOGA GACCAGOCCO CTCCCCTGAT

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20 30 25 35 7 5 25 6 55 80 8 CCAATAACTC CATGGGCTCT GGGACCCTAC CCCTTCCAAC CTTCCCTGCT TCTGAGACTT AGGCTCCGTG CAGCCCTTGG GAACAGTGAG AGGTTGAAGG TCATAACGAG AGTGGGAACT TOCTTOCTTA TOCCACCTGA TOCCAGTOTG AAGGTOTOTT AGCAACTGGA GATACAAAGC ATARANTATC CITICITICM TARARARAR AGRARARAR AGGOGGGGGGGG CTOTGACCCA TIGGIGITCT CIGIATCGIG ATCIATCCIC AACAACAACA GAAAAAAGGA CATCCTTTGC GOCANTAGTT GAAGGACTEC TGTTCCGTTG GGGCCAGCAC ACCGGGATGG ATGGAGGGAG CARTCTACAG CCCAGCTCAT CCAGATGCAG ACTACAGTCC CTGCAATTGG GTCTCTGGCA ATTCCAGGCC TTCCCAGGGG AAGGAGCTGG GCGACCOTCG TTTGAGTCGT CGCTGCCGCT GCCGCTGCCA CTGCCACTGC CACCTCGCGG CAACCCAGAT CCCGCCCCCC CTGTCCTCTG TGTTCCCGCG GAAACCAACC AAACCGTGCG AGCAGAGGCC TITGCTICTC TGCCTACGTC CCCTTAGATG GGCAGCAGAG GCAACTCCCG (2) INFORMATION FOR SEQ ID NO: 291: CAGTOTTOCT GARAGGANAG ANGAGACGAG ANGCTOTTTG CATCOTCCTT TCTGATGATA TICICAAACA GAAGAACCGI CCCAAICGGI TAAITGIIGA IGAAGCCAIC AAIGAGGACA GAGAGGCGCG AGICOGICOC TIOCCACCOC ICGIAGCCGI TACCCGCGGG CCGCCACAGC CGCCGGCCGG GCCGCTTGCG ATCAGGAGCC ACAGTGTGGT ŝ E TCTCCCTCTC GGTCGTCAGA GCCGTCAGCC AGGTCGGGTTG GAGACTCAGC CAGGGCTTCT ACTOTOCCCC TOGGGAATGT GICCCCTGCA TATCTTCTCA CITICOACGAG GAGIICCCCAT CIGCCCCGCC CCITICACAGA GCGCCCGGGG TOAGCCCAGC GITGACGTCA GGCAGGCTAT GCCCTTCCGT GGTTAATTTC SEQUENCE CHARACTERISTICS: GICCTICICC CAGCCCAAGA TOGATGAAIT GCAGTICITC CGAGGIGACA COCCATOGOT TOTOGAGOOG ATTOAAAAGG TGATGACCTA TOAACAGOOA GETTTOTICGE COCTGCTCGE CHACCGCCTG GAAGAGCCGA GECCCGGCCC ACCETTETTE GCCCGACGCC TCGCTGCCGG TGGGAGGAAG CGAGAGGGAA SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 3276 base pairs 291 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320 1260 1910 240 180 120 480 420 360 300 60

မ 23 20 2 35 9 45 6 SS 50 8 GIGGGAIGCG IGCIGIGGAG TICAAAGIGG IGGAAACAGA ICCIAGCCCI TATIGCAIIG TAGGGGATOT CATCAGCATC CAGCCATGCC CTGATGTGAA GTAGGGCAAA CGTATCCATG CTTOTTCTGA TGAGAAGATT CGGATGAATA GAGTTGTTCG GAATAACCTT CGTGTACGCC GAGCTOTAGC ANATCAGACT GGAGCCTTCT TCTTCTTGAT CAATGGTCCT GAGATCATGA TARAGGREAT GOTGERACTE CCCCTGRARC ATCCTGCCCT CTTTRAGGCA ATTGGTGTGR TIGCTCCAGA CACAGTGATC CACTGCGAAG GGGAGCCTAT CAAACGAGAG GATGAGGAAG TOCTOCCCAT TGATGACACA GTGGAAGGCA TTACTGGTAA TCTCTTCGAG GTATACCTTI CICTACGOCG ATTIOGICGC TITGACAGGG AGGTAGATAT TGGAATICCT GATGCTACAG GCAAAITIOGC TOGTGAGTCT GAGAGCAACC TTCGTAAAGC CTTTGAGGAG GCTGAGAAGA AGCCTCCTAG AGGAATCCTG CTTTACGGAC CTCCTGGAAC AGGAAAGACC CTGATTGCTC AGICCTIGAA IGAAGIAGGG IAIGAIGACA IIGGIGGCIG CAGGAAGCAG CIAGCICAG AGCCGTACTT CCTGGAAGCG TATCGACCCA TCCGGAAAGG AGACATTTTT GAGCCAGAGT AACCCATCAG CACTOCOGGA AACCOTGGTA GAGGTGCCAC AGGTAACCTG CATTGATICC GAGGICATGA ACTICICIAGO AGITACIATO GATGACTICO GGIGGGCCIT TOGANCAGTA GCCAATGAGA CTCACGGGCA TGTGGGTGCT GACTTAGCAG CCCTGTGCTC GACGCTTAGA GATTCTTCAG ATCCATACCA AGAACATGAA GCTGGCAGAT GATGTGGACC AGCAGAGGGC ACATOTGATT GTTATGGCAG CAACCAACAG ACCCAACAGC ATTGACCCAG CTCATOGCGA GGTGGAGCGG CGCATTGTAT CACAGTTGTT GACCCTCATG GATGGCCTAA ATGCTCCTGC CATCATCTTC ATTGATGAGC TAGATGCCAT CGCTCCCAAA AGAGAGAAAA GGAAGACATC GGGGGCCTAG AGGATGTCAA ACGTGAGCTA CAGGAGCTGG TCCAGTATCC AGAGOCTOCT CTOCAAGCCA TCCGCAAGAA GATOGATCTC ATTGACCTAG AGGATGAGAC GECCAATETC AGAGAAATCT TIGACAAGGC CCGCCAAGCT GCCCCCTGTG TGCTATTCTT CTANOGACCT CCTOGCTOTO GGAAAACTTT GTTOGCCAAA GCCATTOCTA ATGAATOCCA CCTCAAGGCT AACCTGCGCA AGTCCCCAGT TGCCAAGGAT GTGGACTTGG AGTTCCTGGC TGATGAGCTG GATTCGATTG CCAAGGCTCG TGGAGGTAAC ATTGGAGATG GTGGTGGGGC GOCCAACTIC ATCICCATCA AGGGICCTGA GCTGCTCACC ATGIGGTITG GOGAGTCTGA IGNOGAGCAC CCAGACAAAT TCCTGAAGTT TGGCATGACA CCTTCCAAGG GAGTTCTGTT IGCTGACCGA GTCATCAACC AGATCCTGAC AGAAATOGAT GGCATGTCCA CAAAAAAAAAA IOSCOGICTI GAICAGCICA ICTACATOCO ACTICOIGAT GAGAAGIOCO GIGTIOCOAT ATTOGCCCTA CCAACCGGCC TGACATCATT GATCCTGCCA CITOICCOIG TCCTCAGACC 1560 1500 1440 1380 1320 1260 1200 1140 1080 1020 1680 1620 2220 2160 2100 2040 1980 1920 1860 1800 1740 960 90 840 780 720 660 600 540 2280

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2340 2400 2460 2520 2580 2640 2700 2760 2820 2880 2940 3000 3060 3120 3180 3240 3276

TAMANTGACT ANTGGCTTCT CTGGAGCTGA CCTGACAGAG ATTTGCCAGC GTGCTTGCAA GCTGGCCATC CGTGAATCCA TCGAGAGTGA GATTAGGCGA GAACGAGAGA GGCAGACAAA CCCATCAGCC ATGGAGGTAG AAGAGGATGA TCCAGTGCCT GAGATCCGTC GAGATCACTT IGAAGAAGCC ATGCGCTTTG CGCGCCGTTC TGTCAGTGAC AATGACATTC GGAAGTATGA GATGTTTGCC CAGACCCTTC AGCAGAGTCG GOGCTTTGGC AGCTTCAGAT TCCCTTCAGG GAACCAGGOT GGAGCTOSCC CCAGTCAGGG CAGTGGAGGC GGCACAGGTG GCAGTGTATA

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TOGOCTOCCT GAACTITOTT COCTGGGGT GGGGCCCTT CCCCAGGAGA GGGACCAGGG

STOCOCCCAC ACCTGCTCC ATTCTCCAGT CTGAACAGTT CAGCTACAGT CTGACTCTGG

CACAGAAGAC AATGATGATG ACCTGTATGG CTAAGTGGTG GTGGCCAGCG TGCAGTGAGC

15

ACAGGGGGTT TCTGTTGCAA AAATACAAAA CAAAAGCGAT AAAATAAAAG CGATTTTCAT

20

MOGIAGOCO GAGAGICAAT TACCAACAGO GAATICOGOCC TIGOOCIAIG CCATITICIGI TOTAGITITGS GECAGTICCAG GEGACCTIGTIG TOCOGNISTIGA ACCAAGGCAC TACTGCCACC IGCCACAGIA AAGCATCTGC ACTIGACTCA ATGCTGCCCG AGCCCTCCCT TCCCCCTATC CAACCTGGGT AGGTGGGTAG GGGCCACAGT TGCTGGATGT TTATATAGAG AGTAGGTTGA ITITATITITAC ATGCTTTTGA GITAATGTTG GAAAACTAAT CACAAGCAGT ITCTAAACCA raaaatgaca tottotaaaa ggacaataaa cottoooton aaatgoomra aaaaaaaaa

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420 480 540 009 9 720 ဝ

S 120 180 240 300 360

PIGCAATIGGT TGAATTCCCC TCCTCACGCC AGCCTAGGAG AAGAAGTTCG TAGTCCCAGA GGTGAGGCAG GAGGCGGCAG TTTCTGGCGG GTGAGGGCGG AGCTGAAGTG ACAGCGGAGG

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SEQUENCE DESCRIPTION: SEQ ID NO: 292;

(X

LENCTH: 1695 base pairs

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(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 292:

6

TYPE: nucleic acid STRANDEDNESS: double

TOPOLOGY: linear

5

AAAAAAGGG GGCCCTCTA AAGNNCCANN CTTCGT

35

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25

CCTTORCCOG AGGITCOGGG ACCOCTICGG CTGAAGCATT TGACTCGGTC TTGGGTGATA

8

TCCGAGAAAA CAGCAACAAG CTGAGCTGCT GTGACAGAGG GGAACAAGAT GGCGGCGCG AAGGAAGCCT CTGGGTGAGG ACCCAACTGG GGCTCCCGCC GCTGCTGCTG CTGACCATGG

25

COGAAGCAAC GGTCGGTGGG GCGGAGAAGG GGGCTGGCCC CAGGAGGAGG AGGAAACCCT

မ 23 8 5 6 35 5 55 S 25 S CTOGTOGTOG GGGCGGCCG CATCGGCTGC GAGCTCCTCA AGAATCTCGT GCTCACCGGT TGANTOTOGT TOOGTTOGAC TOGGCTGAAG TACAAAGTCA AGGAGAAGAA ACGAATGCAT CAGATCAAGA ANGGANTGOG CTANATICANC TGGNTNTGAT CCAGTITURAA CTTTTTTACCA AGCTTTTTAA GAAGCCGAAG CCAGAGCTAG AGCATCTAAT GAAGATGGTG ACATTAAACG TATTTCTACT GATGCTGATC AAGAAGTATC TCCTGACAGA GCTGACCCTG AAGCTGCCTG GGAACCAACG GAACCTATAC ATTOCATOOT TICOGCAAAG TACTICTICA ACCACTICTI TOGGGAAGAA ACAGCTGGGT ATCTTGGACA AGTAACTACT ATCAAAAAGG GTGTGACCGA GTGTTATGAG GCCCGAAACC ATGITAATAG AATGIGCCTG GCAGCTGATG TTCCTCTTAT TGAAAGTGGA TATAATOTOG AATTITITOCO ACAGITIATA CIOGITATGA ATOCITIAGA TAACAGAGCI CTGCAGTTTT ACCCGAAAGC TAATATCGTT GCCTACCATG ACAGCATCAT GAACCCTGAC TTTTTGTTTC AAAAGAAACA TGTTGGAAGA TCAAAGGCAC AGGTTGCCAA GGAAAGTGTA TICICCCACA ICGACCIGAT TGATCIOGAT ACTATIGATG TAAGCAACCI CAACAGACAG ANDGEACTOT COCOGGOET GEECECOGAAG CTGGETGAGG COOTOGECGG GGGEEGGOTR TOTTTTTTCA AAGAGCATOG AGACTTTGAG AGTTCATTTA GCAGAAAAGG GGGATGGAGG GAATGAACCC CAGTTAGGCC TGAAAGACCA GCAGGTTCTA GATGTAAAGA GCTATGCACG TOTCATCCTA AGCCGACCCA GAGAACCTTT CCTGGCTGTA CAATTCGTAA CACACCTTCA ACAACCAAAC CCAAGAAAGA AGCTTCTTGT GCCTTGTGCA CTGGATCCTC CCAACCCCAA GGAAGGATIG AAGATITIAT CAGGAAAAAT AGACCAGIGC AGAACAATII TITIGAATAA AGGGAACATT ATTECTOCTA TIGCTACTAC TAATGCAGTA ATTGCTGGGT TGATAGTATI CCTCAGGATG CATATTTTCA GTATGAATAT GAAGAGTAGA TTTGATATCA AATCAATGG TGAGCTCATA TOGGATAAGG ATGACCCATC TGCAATGGAT TITGTCACCT CTGCTGCAAA AGATGACATO AGOTATOTOT TGACAATGGA CAAACTATOG OGGAAAAAGGA AACOTOCAKT TGACTICCIC CAGGACIATA CITTATIGAT CAACATCCIT CATAGIGAAG TANTANTCAC AAGAAGTTGT CAGAATTTGG AATTAGAAAT GSCAGCCGGC CCAAATTGAA GATGGGAAAG GAACAATCCT AATATCTTCC GAAGAGGGAG TGTTCTCACC TTACAAGACA AGATAGTGAA AGAAAAATTT GCTATGGTAG CACCAGATGT TIGITAIGTA IGIGCCAGCA AGCCAGAGGT GACTGTGCGG CTGAATGTGC ATAAAGTGAC CCCGGGTGCN GATTGGCAGN GCCTCCGCCG CGGCTCGTGG TTGTCCCGCC ACCTAGGAAA TTCAAGCAG AGACOGAAGO 1140 1080 1020 1680 1440 1380 1320 1260 1200 180 1620 1560 1500 360 300 240 120

960 900 840 780 720 660 600 540 480 420

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ

E NO:

8

GWICICCOGG TTATTTICCAG TOGGTGTAAA AGCAGAGCTG GGCCTTTCCC 180 120

50 AATAGTTGTA AGTTTGCATG CATGATGGAA AAATAAAAAC CTGTATCTCT GTTAAAAAA

45 TACTTOGTAA TITGACACCC TGTTAATAAC GCAATTATTT CTGTGTTCTT AAACAGTATA

1500

1501

1440

6 GARATTOTTO CTOTOTTCTO TGARARTARC CTCCCCARAR TARTTAGTRA CTGGTTGTTC CATTAGCATT GITATCAGCT TGTACTGGTC TCATAACTCT GGTTTTGGAA GAATAATTTG TTGAGACCCT TCACCAGAAT GTCAATCITT TITTCTGTGT AACATGGAAA CITGTGTGAC 1380 1320

AACWAGAGAG AGACTGITCT GITGIAAAAC TCITTCAAAA ATTCIGATAT GGTAAGGTAC 1260 1200

ઝ ATTAACAAGG AGTCTCAAAA AGAAATGAGA GGGATGCTTC CTTTNCCCTT GCATCTACAA TTTCCTTTTA ATGTTAACTA ATGAAGTTCC AGAGATGOGC CTTAGAAATG TOTTTTAAGA 1140

30 TAGTETETT TGATGGTGAT AGTGATGGGG TGCACTATCA TAATCACATE AGGTETGETT 1080 1020 960

TIGITIATIG AAAAICCAAG ACACTAIGCC AAIGCAACCG IGACIACITI GGGAGAIIGG

GCATCACTTA TCAGCTATOG TCAACCTGGT TTCATCTGTA TCTCTCTCTT TTCACCTGTA TCAGACATOC ACAGAAGTOG AGAGGATGOT CCTTGGACCC MATGTGTCCA TCACCTAGCT

900

25 AGICIGCITT TIIGITIITIT GITAITAITT TITITITITT GCICIGIGIT AIGGACAITT 840 780

TIGCCITIAA AGICTIACGA CITICCCCAT TITAGICTAA IGGGAAGATA CAGAIGIGCA

720

8 TAGTAGATTT TATAAGCCAC AGAGACAAAC CAGAAACGGA ATAATGTTAC TITGGATGCT TTATTTTTT CITICIAGGIG IGCCITIGIA CAIGCAGAAG AAIGCIAIAI GCIGCACAIT 660 600

. 15 TTATGAACAA ATAGGATOCC TAGTTGAGGA TGTTCCCAAA GTTTTGTCCA ATCTTATCAT TOTOTOAGIG ATAGAGTATG GGAGGGACOT COCTAGOTIG GAAAATGAGA ATTGAAGGG

540

0 TOCACTOCOC CTOCAAAATC AGACAGAAAT GGCTTGAGAA GCCGCAGOGG AGCATGCCTG 480 420

CCAAGGGTGT GCTAGGTGGG GGATTTGGAG CAAAACCGTC GAGTAGGCAT GATACTGGTA

360

ICITITICITA GOGAGGCCIC ATTOTANGTT CCTCAAGAGA GTCCTTGGCT TAAAGCTGTA CIGAGGGIGG GTAAGAAGGA CIGTATCTAC ACCIGITCIT CCCTACCTIC 300 240

TCTCTTATCC ATCTGTTTTG

S

TAGTOTOCAA TATOGAGCAT CTOAAGCTTO TOOTGGGGA TGGGGATTOG GATOGGCAGA

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WO 98/39448

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SEQUENCE CHARACTERISTICS:

E

55

(2)

INFORMATION FOR SEQ ID NO:

294:

(A) LENGTH: 2683 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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GGACOTTGAA TTTGAAGTTG TTGGTGATGC CCCGGAAAAA GTGGGGSCCA AACAAGCTGA	1740	THE PROPERTY OF THE PROPERTY O
AGATOCTOCC AAAAGCATAA CCAATGGGCA GTGATGATGG AGCTCAGCCC TCCACCTCCA	1800	1000/10CTGC AGGGGCCCCC TGCGCCCCAA' GALACICCIG: CICACCAGG, CCGAGGGCTGC
CAGCTCAAGA GCAAGATGAC GTTCTCATAG TTGATTOGGA TGAAGAAGAT TCTTCAAATA	1860 5	
ATOCCGACOT CATGAAGAAG AGAGAAGCCG: CAAGAGGAAA TTAGATGAGA AAGAGAATCT	1920	GAGAGACAGG TACCGGCTGG ACGATGGCCG CCGCGTCCGG GACCTGGACC GAGTGCTCAT
CACTGCAAAG AGGTCACGTA TAGAACAGAA GGAAGAGCTT GATGATGTCA TAGCATTAGA	1980	
TIGNACAGAA ATGCCTCTAA ACAGAACCCT CTTACTATTT AGTTTATCTG GGCAGAACCA	2040	
CAMPATTANG TOCHTHISTIC CAAAGGGAAA AAATTIGAGAG CAGTIGACTIG AAAAGGATTICA	. 2100	AGCLAGGCC GTGAAGTCCA GTGGCAGGTG TTTGTCCCCA GTGCTGAGAG CAGAGAAAA
INCLUSIONAL SANCTARING SANTERINGS CONTRACTOR	V - C - C - C - C - C - C - C - C - C -	CTCATCTCGC TGTTGGCTCG CCAGTGGGAG GCCCTGTGTG GCCTGAGCTG CCTGTCGAGC
ותיותניון השתתחור שווותיוש שהיחוושה ההיוות שותיומים		TCACCEGGETA GCCCAGGCCA CAGCCAGCCT GTCGTGTCCA GCCTGAGGCC TACTGGGGCA
TOWARDINGS AND THE TANDARCOUT TOWARDAND TOTAL ACCURICATE	0777	GGGCAGCAGG CTTTTGTGTT CTCTAAAAT GTTTTATCCT CCCTTTGGTA CCTTAATTTG
AGATAAAACA ACACAATGCA TOTTOCCITT TIAALGIAAA TACCCITAGS TATCATTAAT	2280 20	ACTIGECTICS CAGADATICTS AACATGTICTG TGTICTTGTS TAATTICTTTC TCATGTTGGG
agiticaaaa tattsingsii tagtaaagit gataccinssi tataaatatt argectitat	2340	AGRAGAMIG COGGCCCCT CAGGCTGTT CGGTGTROCTG TCAGCCTCCC ACAGGTGGTA
TITITGCTAG AGAAGAATT ATTITITAGCC TAGATCTAAC CATTITICATA CICTIAACTG		CAGCOGIGCA CACCAGIGIC GIGICIGCIG INGIGGGACC GIIGITAACA CGIGACACTG
attgaaacag attcaaagaa gtatcgagtg ctatgcattg aaacttgttt ttaaatgtta	2460 25	TOGGRETICAL TITYTCTTCT ACACGICCTT TCCTGAAGTG TCGAGTCCAG TCCTTTGTTG
GATGGCACTA TOTATATAA TOTAAAACAA TOTTAATTTA CTCAAGTTTT CAGTTTOTAC	2520	SATESTIMOSTA STEPS TO THE STATE OF STATESTIMOSTATES AND ASSESSED ASSESSED.
OCCUGINA GICUGIGIAA GAACCAAIT ITHOIGIATI GITACAGIIT CAGGITAITI	2580	
atatticgato tittigiaaa ctcaaataac gactatactit atggaccaa taaatggcay	30 2640	_
and and and and and and and and and and	2681	TCCAGACAAG TGGGAGACC CGTGGGGGCA GGGGACCTGG AGCTGCCAGC ACCAAGGGTG
	35	ATTOCTOCTO COTOTATION CTATICOLAT AAAGCAGAOT TIGACACCON MAAAAAAAA
	1	AAAAAAAA AACH
(2) informaticm for SEQ id no: 295:		
(i) SEQUENCE CHARACTERISTICS:	40	
(A) LENTH: 1454 base pairs		(2) INFORMATION FOR SEQ ID NO: 296:
(B) TYPE: nucleic acid		
(C) STRANDEDNESS: double		(i) SEQUENCE CHARACTERISTICS:
(b) 10k0,00x: 1,10ear	45	(B)
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 295:		0 8
GGACTCGGGG TGGCTCTAAG GGGCAGGGAT AGGGCTGGGG AGCGCCGGCC TGTGGCCCTG	09	Training 110000001 (7)
ACCAGCCCCT TCTCGTGCAG GTTCCACCC GATGCAGGTG GTCACGTGCT TGAGGGGGA	120 50	
		ACCCTGGCAT GCCCCACAAA CAGATCACCA GCCAGCTTAC ACAGGCATTA ACTCTCCTCA
CAGCTACCTG ACCACTGCT TOCTCCAGCA CCTCATGGTC GTGCTGTCCT CTCTGGAAGG	180	ATGAGGAAGA ATCATTCACA ACTGAGCAAG ACATTCATAT GATCATTFAA GGAAGTGTTT
CACGOCCTICG CCGGAGCCTG TTGACAAGGA CTTCTACTCC GAGTTTGGGA ACAAGACCAC	240	
		CCCTTATIGIG TTAGCAAGTA TAATCGGCTA ACTCCTAAAT CCCAATGAAT AGTCCTAGGC

 TGGACAGCAA TGGGCTGCAA TTAGGCAGAT AAAGACATCA GTCCCAGTAA ATGAATCCAT AGACTICATICT AGCACCAACT ACCATTAGCA CTATGTTAGG AGCTGCAAGG CCCCAAAGTA

AGGGAAGATG GAGAACTACG AGCTGATCCA CTCTAGTCGC GTCAAGTTTA CCTACCCAG TCAGGAGGAG ATTGGGGACC TGAGGTTCAC TGTGGCCCAA AAGATGGCTG AGCCAGAGAA GOCCOCAGOC CTCAGCATOC TOCTGTAGGT GCAGGCCTTC CAGGTGGGCA TGCCACCCCC

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GAAGATGTGC ATAATGTCTG CTCTTGTGTA GCTCAGGAGA CAATTCCAGC ACAGACACTA

40 ઝ 30 3 8 55 8 GAGAACAAGG CCAACCAAGG GACAGACTIG AAAGCACTTA GATGTTTAAG GAGGAGAAAG GATTOGRAGICA ACCOCCAAGGA TATGCAAGAA GGOCATGATG AACCCCCTTC CCTCTGGCAG GCTABACTGC TCTTAGCTTA GAATTATGCT TTACTAGAGA TCTAGCAGAT AAGTGGGTTA ANATOTAGAC ANTAGAAGTO ATOGATAGCA GOTTTTCCTC ANATOTOTGA CTCCTCAGGG TTTTTTCTTT GAGAAAGAAG TOGACTGGGG CACAACTTTT AGTCTGAGGG GAGCTAGTGG AACAAAACCT ACTATCACCA TCATCCTTCA ACAGCCACAG TCTGAATTGA GCCAACATTT TCAATTICCA TTAACICAGA ICAGCCATIG IGATICACCA TIIGICAGGC ICICAGGTII CCTCTAGCGC AGTAACATTT GCAGAATTGC AGATTTTCCC CCAGATACTA GGAGGAAAGG GODANGCITT GACCAGICCT TECCTITIGC CAAGIICAGC CAGIICICCG CIGCIIGCAA ATCACTACCA TOOTGEACT AGTEATATAG CTTOCAGACA TGAGGGAGAC ATCAAACAGG COCTOCOTTY COTTYTOOTA TOTACTICCY TOATACTIGG TITACTICATIC ACCORDIGADA CATTOTOTOT TOCATAAAAC ACTATATTTT TITOGAAATG TIACIGICCA AAAGCCICTI THITTATTIT GICATOCICI TGAAAATGIT IGACCATIIG TAGIATACAC AGIGAAACTI TOTTTTAAGA TAGGAAAAAA AAATAGTGGG CAAGGTGAAC ATCAGAGGTA AATTTGTGTG X GGTGGGGAAG SEQUENCE DESCRIPTION: SEQ ID NO: 297: GOGICGIGGT GITTIAAAAG CATAAGITAC CIGITIGCAC 180 120 240 420 360 300 660 600 540 480 6 720 900 840 780

3 INFORMATION FOR SEQ ID NO: 297: 20

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ATCAGGGATC TTTTCACACT GCTGTTTTTT CCTCTTTGGT CCTTCTATCA CTAAAACTCA

TCTCATTCAG CCTTACAGCA TAACTAATTA TITGTITITCC TCACTACA

828 780 720 660 600 540 480 420 360

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OCCADACTOT CCCAGCCAGA TOTOTOCCCC CACCCCATOT CCATTTACAT CTCCTTCAAT GIGIATICAG AGCACCTYTC CAGAIGCACC ATGCATGCIC ACAGICCCIT GCCIAIGIGI

5

CAGTGAGGAG CCCAGGGATG GAGGGGGTTC CTGAAGTATT GCAGTTGGCT GTAGTAGCTG

AAATGAGGOT AACAGACCAG ACTGCAGCAA GTTATCAGAT TCCTCAATCA GATGCACTAG

AAGCCAGCOT GTGCCCTCTG GTTTAGTGAG TGTAATAGAG TCCCTGGCAC

TACCCATCCA AGACCTAGAG CATGAAACAG GGCCCTTTCC AAGTAGGCTC TGGGTGTCCT

960

AGTICITITIC CAUGITACCG ANACIGIAGC CAGITACAGI TIACICAGGA ANACGGIAGA

TCAATTCAGC CATOGTAGTS CTGGTTGGCA GGGATTGGTA ACGGAGAGAA CTGCTCATCA

GCCCACCTCA AAAGGYACYT CTTCTGTAAA GCTTTCCCTK GGTATCAGGA ATCAAAATTA

S

ACCCCCAGG CCTGAAGCTG GCCCTTGAAG GATGGATGAA ATTTGGATAG AGAATGAGGA CAGTTAACGC TGAACTGCAG CTGCAAGTAA TAGCAWGAAC AGTCAGAAAA ATACCTTATG

AGACAGAGG NCTCCAAGTG AGAGAAGCAT GAAAAATGAG CARGGGCCTG GATCAGTGGG

Ξ SEQUENCE CHARACTERISTICS:

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OCCAMANCIC ANGCCTIOCC TITINGGAGG CCACCAGCAG AGGGACTIOG TCCTCCTTGT

CTGGTACTTG TGTACATGCC GGTGACCTGA GGACTCCACT CACACTGGCG AGCAAAAAAGG

GAGCAGIGAT TOTOTTTOT CICODOACCO CONGCCTIT GITACCAACA CCAGITITCCC

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(A) LENGTH: 2416 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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GAGTETETGA GETTTTCAGT CCAAATETTT GCAAGGETCA AAATGCCACA GAACETETEC GOCTOCAAAR COAGAAGAGA GCCGGTOGAG TGTACTTGTC CCTGACAGGC TGACCTACCT TOTOTTATAT TCTOCTTOTO AATAGCTOGA GCAAACCTOG GOCTGACACG CGTAAGSTAG TTATATATIG AGIGAIGAAT IGAICCICIT TITICCCTAA GGGATAIGAA TIGITITITCI GIATTITICI GIICACAGIA TIIGIGIGIG IGCIIGITII GGCAGCICAI TIIGGCIGIA TGATAAGTGC TTTAAGCAAT GTCCATACCC CGTCAAGACT CCCAGCTTAG TCATTTTCTT

1740

1680 1620 1560 1500 1440 1380 1320 1260 1200 1140 1080 1020

TETTETECCE ACTECCEATG GEAGGGACCG GACCATECET ACATGGAACA TGCTGTTCCT

1920 1860 1800

6 3 GAATIGIGAA GGICGGAGIA GITAGAICIT TAGCITTTAT TCCITATITT TITGIAITAC CATCCCAGGG CTITOTGACA GICTCTAATI CCCTICCCTI CICGITAAGA AICATATIOT CTICCINGA CACCCGAGCT GCTIGCCCAG GGICCIGTIT CCCIGCINAC ICCAGAGANG GATGCITICTG TAATCGACCA CCTAGCCTTC TCTCTCCCCT CCCGTCCTCC CCCAGAATCA CCAGCCCCTC CCATTGCCAT GGCAMAACAG GTACCTTTGG GGCATGGGGG CATTACATGG ATAGTAGCTT TCAGACCATA CAGTATTCAT TOGGTTACTC CTATTATTAT CAAGTAGCTG

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2160 2100

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GAGCACTICCC TCAGCCTTTG

CACATGGTAA TGAAGCACTG TTTTTAAATA AAAGRGRGAA

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TCTCCATGTG TATAAAITAT TGATCATGTT GCTGGCTTTT ATAAACTCTA AGCGAAGGAG

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2 INFORMATION FOR SEQ ID NO: 298:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 545 base pairs

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(B) TYPE: nucleic acid		
(C) STRANDEDNESS: double		TGGAGACCAA GTGGAGGAAC GGGGACATCG TCCAGCCAGT CCTCAACCCA GAGCCGAACA
(D) TOPOLOGY: Linear		CTGTCAGCTA CAGCCAGTCC AGCTTGATCC ACCTGGTGGG GCCTTCAGAC TGCACCCTGC
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 298:	vs	ACCICATIVOT CCACCGAGGT GTCACCATGA ACCICATGGA TGAGGTCGCC GGGATCGTGG
GAATTCOGCA CGAGCCATGC YTGGCCTCTC CTTGATTCTT ACAGTCACTT TGTTGGCTGT	. 09	CTGCACGCCA CTGCAAGACC AACATCGTCA CAGCTTCCOT GGACGCCATT AATTTTTCAÑG
TICTGACTCA GCACCTACCT GCATTGTGGC CAAAGGATGA CCTATTCCTT CTCAGGAGGG	120 10	ACAAGATCAG AAAAGGCTGC GTCATCACCA TCTCGGGACG CATGACCTTC ACGAGCAANA
CAAAAAIGIG GAATAGIGIC IGICCAIGCC ICTCCTCAIG GGCTACCACC ICTGCCACCG	180	ASTOCATIGA GATCGAGGTG TTGGTGGACG CCGACCCTGT TGTGGACAGC TCTCAGAAGC
TOSTTANTCA STANCANCCA GOAGAGARGC TOCTGGAACT GACCTCTGGG AACTCCCTGG	240	GCTACCOSSC COCCAGTOCC TTCTTCACCT ACGTOTCOCT GAGCCAGGAA GGCAGGTCGC
ATGGTTTGGT GCAGGAATGT AGTAGGCATA CACGTGGTTG CGTGGATCTG GGCCCTCCTG	300 15	TOCCTOTOCC COAGCTOOTG OCCORAGICCO AGGACGAGAA GAAGCGCTTT CAGGAAGGCA
ATGTGAGTAG AGAGGTAAAA GGSCACCATC TCCTTGACCT YTGGGGAACT CATCCACAAA	360	AAGGGCGTA CTTCAAATG AAGCCAAAGC GACAGGCCA CGCGGAGCCT CAGCCCTAAA
GAAGATGTTT CCAAGATGCT TCTGAAGATT GSCTAAAAAT AGCOGGTTTC CACCCCCTG	420 20	CITCCTICCT CTGCCACTGG TGCCTCGAGT AGCCATGGCA AGGGGCCCAG TGTCCAGTCA
AATGCATCCA TTCTAGAATG CTCCTTCACC AGGACCAGAG AACTGATTTA CAGAAGTGAC	480	CTTAGAACTT CCCCCCTTGG CCAAAAACCC AATTCACATT GAGAGCTGGT GTTCTCTGAA
ATGAAAACAT TOCATOOCAG AATTIGGANT ACCTOAAATT NAATTICIAC CTATTAAAAA	540	GITTICGIAT CACAGIGITIA ACCIGIACIC ICICCIGCAA ACCIACACAC CAAAGCITITA
NAAAA	545 25	TITARATCAT TOCAGIANCA ANGCIACACA GIGTIGNOCO GAGGGGGGGG AGGUSTIGGS
		CAGAAACCCT COGGAATOCT TCCGAGCACG CTGTAGGSTA TGGGAAGAAC CCAGCACCAC
(2) INFORMATION FOR SEQ ID NO: 299:	30	THATAAAGCT GNIGCTIGGC TGGGGAAGNA
(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1510 base pairs		
(B) TYPE: nucleic acid (C) STRANDEINESS: double	35	(2) INFORMATION FOR SEQ ID NO: 300:
(D) TOPOLOGY: linear		(1) SEQUENCE CHARACTERISTICS:
(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 299:		(A) LENGTH: 997 base pairs (B) TYPE: nucleic acid
GGCTCTGCTG GGCATCATAC TIGTCACTOG GTAAACAGTT TGCCCACTTA CCGCAGATGA	60 40	STRAN
AGCTOCTITICS CAGGGCTCTC CAGCTCTGTG AGTITGGGAG GCAGGCATCT TCCAGGAGGC	120	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 300:
TRATESCRISC CCARGGART GROSSCCC GOCGAGGGTG CTGCGCTCCC GTCCAGGTGG	180 45	AGGIAGIGAG AGACACATIA CACCIAACCA ACAAGAAGAA GGAICCICCC CCTIAITAATT
TTGGGCCCAG GGCTGATCTC CCACCTGTG GAGCCTGCAT TACTGGAAGG ATCATGGGCC	240	TAACTATGTT TACAGGGAAT GCGTACATTG TGGCTTCCCG AGNATTTCGT CCAACATGT
CAGATGATIC CAACGTOOCC GOCAATGTOC ACGOCGOGAC CATCCTGAAG ATGATCGAGG	300	TIGAGARC CINANICCA ACARCIGNIT GANIGGGIAA AAGACACTIA TAGCCCAGAT
AGGENGGOGC CATCATCAGC ACCCGGCATT GCAACAGCCA GAACGGGGAG CGCTGTGTGG	360	GAACACCTCT GGGCCACCCT TCAGCTGCA CGGTGGATGC CTGGCTCTGT TCCCAACCAC
COGCCTIGG TOGTSTCGAG COCACCBACT TCCTGTCTCC CATGTGCATC GATCAGGTGG	420	CCCANGTINGS ACATETICAS ACATISACTIC TATTGOCAGG CTGGTCANGT GGCAGGGTCA
COCATOTCAG COCOGAGATO ACCTACACOT COAGGACTO TOTOGAGATO CAGOTOAACO	480 55	TGAGGGAGAC ATCGATAAGG GTGCTCCTTA TGCTCCCTGC TCTGGAATCC ACCAGCGGGC
TOATISTICCOA AAACATOCTIC ACAGGTGCCA AAAAGCTGAC CAATAAGGCC ACCCTGTGGT	540	TATCTGCOTT TATGGGGCTG GGGACTTGAA TTGGATGCTT CAAAACATC ACCTGTTGGC
ATOTOCCCCT GTCGCTGAAG AATOTOGACA AGGTCCTCGA GGTGCCTCCT GTTGTGTATT	009	CAACAAGTIT GACCCAAAGG TAGATGATAA TGCTCTTCAG TGCTTAGAAG AATACCTACG
CCCGRANGS GARGARGES GARGARITAS AGRICANAS AGRICADAS MINITARAS	09	

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GGAGCTATTG AAAAGGCTGT AATTTGGCCT CAGTATGTCA AGGATAGAAT TCATTCCACC GOGGETOCTO TIGGICTIGG AGENTICITOC TACTATOSCT ISOGACIGIC TAATGAGATT GAACCATCGA TOGAAAAAAT ATTTAAAATT GATCAGATOG GAAGATOGTT TOTTOCTOGA AAAACAAGAA TIOGGATCCG GCGTGGGAGA ACTGGCCAAG AACTCAAAGA GGCAGCAFTG GTGAAGAATT CCATCACGAA GAATCAATGG CTGTTAACAC CTAGCAGGGA ATATGCCACC TOTCTCCGGA CACTACCTIC TAGGGITTIC CACCCAGCIT TCACCAAGGC CICCCCTGIT CATTICAGAT CIGCICGGTA GACCIGGTGC ACCACCACCA TGTIGGCTGC AAGGCTGGTG CCAAAGCATC, TIGCTIGGIT GCTACATICI GGIGIGATGG GIGCAGIGGI GGCICCICIG ACCCCTOTIC TCATGAACIT CATGATGAGA GGCTCTTGGG TGACAATTGG TGTGACCTTT TATATOTACT TAGCAGGGAG TATTGGTTTA ACAGCTTTGT CTGCCATAGC AATCAGCAGA ттанвоссан состиновае совальский истосанове амистансе возвиссана CHOGGAGNOG GCCTOGÓTICT COTCTITIONG TCCTCATTOG GATCTATOTT TCTTCCACCT GOCCICICCA CIGIGOCCAI GIGIGOCCC AGIGAAAAGI IICIGAACAI GOGIGCACCC GEAGCCATGG TIGGAGETGG AATGETGGTA CGATCAATAC CATATGACCA GAGCCCAGGC ξ. GGGTCCTCT SEQUENCE DESCRIPTION: SEQ ID NO: 301: TCTCATCAGA GCTGCATGGT ACACAGCTGG CATTGTGGGA 360 300 240 180 120 540 480 420 660 60 900 780 720 60

20 ઝ 30 25 2 5 50 3 8 Ś CARAGTARGG ARARARARA ARAGARARAA ARCTCGA GCCTTTGCAA TTCGTGGCAT CCTTTAGGAT AAGAGGGCTG MTATTAGATT GTGGGTAAGT GGGCAAGAGC TTATAAGGCC ATCTATGGGA CTGAACTTTG AGACACACTA TGAGAGCGTT GCTACCTGTG TIGATOGAAA GAGAACCIIC CCIICIGIAC IOTIAACITA AAAATAAATA GCICCIGATI CCTAGTAGTT 2 TYCCATYCTG TGGAGCTGCC GTTCCTAATA ATTCCAGGTT TGGTAGCGTG GAGGAGAACT AGATCTTTTG INFORMATION FOR SEQ ID NO: 301: Ξ OGGATATOTO OTAGAGCACT TOATTTCAGT TGAATGCCTG CTGGTAGCTT CCTCCACTAA CTTTCTCACT AAGTGAGAAT GAGAACTGCT GTGATAGGGA COTTIGUADAT TOCTIGOCTIGG GTGRATIGOTIG CITIGTTICTICT CACCCOTAAC ATGTACAAAC ATGCTCAGAA CITGCTGGGA CAGTGTGGGT GGGAGACCAG SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2345 base pairs (A) LENGTH: 2345 base paix
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 960 540 997 900 600

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(2) INFORMATION FOR SEQ ID NO: 302:

SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (A) LENGTH: 2369 base pairs

(C) STRANDEDNESS: double

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TOTTACATIC ATCTOCTOAA CITAACAAAA CIGITCATCC IGAAACAGGC ACAGGIGAIG

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TYMAAATTTA GIAGGTICAC IGAGIAACTA AAATTIAGCA AACCIGIGIT IGCATATITT TOTTTTOSTG AATOTGAAAA CTAAAGTTTG TOTCATGAGA ATOTAAGTCT TTTTTCTACT GTAATCCTCT CCCAAATAAG CACACACATT TICAATICTC AIGITIGAGI GATITTAAAA

CAGAGAGAAG GAGTCACCTG CAGTCTTTTG TTTTTTTAAA TACTTAGAAC TTAGCACTTG TITIOGRAFICE AGRATIATION ARTHARISTIC ATRAGEGRATI TOGRAFITITIS GIVARAGOGRAF

TOTTATTGAT TAGTGAGGAG CCAGTAAGAA ACATCTOGGT ATTTGGAAAC AAGTGGTCAT

1680 1620 1560 1500 1440 1380 1320 1260 1200

1740

CATICICCIO CIGITOCIIC ICAGIOCICI CITICCAATA IAGANGIGGI CANGITIGAC TIOTACAGAA TOTTAATCAT ACAGAGAATC CTIGATOGAA TIATATATOT GIOTTTIACT

TTTGAATGTT ACAAAAGGAA ATAACTTTAA AACTATTCTC AAGAGAAAAT ATTCAAAGCA

TCHARTATOT TOCTTTTTICC AGARTACARA CAGTATACTC ATGRITGCTA AGTOTTTTT

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TATTITIOCA TATTIATICA ACIGICIAAT IGAATACAGC TIGCICTIGI CACCICIICA

2040 1980 1920 1860 1800

AGCTITCAAG CCTTTATAGA AAAGCTICTT TOTGGCTTAC ACTGGAAATT ATGAAAGCAG

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TITITICICCI AAGACTITIG GITTCICGCA TIGCCICICA GACTAAGCAC TAAAAAGCAA

AATGCTTCAT TAGTTTOCCC TAGCAGACTT TTACTTCTCT AGCAAAACAG AACTAGTNCT GTCTTAATGA AATATATCAA CCCAAAAGTG TAATGAGGAA

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ATTIGIAAGI CCITIGATAC AGAAGAGITA TATTIAGGAG GNCTITAAIG 2340

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GACTOAGCTT CIGOCTICTO TOCTACATOA AATATCTTOT TIAATOGGGC AGATATGCAT

TARATROTTT GTACAROCAG CTTTCGTTGA AGTTTRAGAAG ATRAGARAACA TGTCATCATR

CCGGTAATGT GATGCCTCAG GTCTGCCTTT TTTTCTGGAG AATAAATGCA

THIOGROTTC AMAMENTGA TECCHITAME TEGRITOCTICA GINTETMENT GGATHERITM AGCHTOTICC TICTOTATGA TACCCAGAAA GTARICAAGC GTOCAGAAGI AICACCAATO ACCACCOTOS CTOSTOCCAC TCTTTACTCA GTOSCAATOT ACGSTOGATT AGJTCTTTTC

AATATATTTA TOCGAGTTOC AACTATOCTG OCAACTGGAG GCAACAGAAA GAAATGAAGT

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PCT/US98/04493

AAT GAGGCCAGC AAGCGCCTGT CCATGCGGTG AGGSTCATTC ACCCATTTGT

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(D) TOPOLOGY: linear		CCAGGTAAAT GAGGOCCAGC AAGGGCCTGT CCATGGGGTG AGGGTCATTC ACCATTTGT
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 302:		CAAGAACGCC TTCCTGTACT TTCTTGATGA GGOGCTGCTT AATGTTGTTA TTGGTGAGG
TITITITIT ITTITITIT TITITINCAG ATCATIGITI ATTIATIACT TCAGATAAA	09	5 GATGTGTTGT CATGTCAAA AGTAGGAAGT TCTGTTTCTC TGTTGTCAAT ACACCCTTTT
AGATAGTATA CATATTAGGG AATCCCTTAA AATTCAACTC TAGAGTTATA CACCATCTAG	120	CCACCAGGIT TITAGCTAAT CGITCCGGIA CATTICITAA CIGAIAAIGC AATHITAAIG
TACTITITICA ATGANIGITA ACAACAACA AAAAANCIC TAAACACTIG AAAGOCCAAC	. 180	GAITCCANGT CTCACCACTA AGTAATTCAA TCCAGITCTG GACGGITTCT GGAGGCJGAA
tattaacatg gactatggta ataaaaatt tigacatita atttgticaa catatagtat	240	TITECTIANE ATSCITCAGA GCITCATCAA GAAGAACATC CCCTGITGGA GCATCTGACT
ttacattatg aaaccaatgg tgatgataca ataaagtgat aaagaaatag taaaaataaa	300	TACAGATTAC CTITCTIGIT AATAGACITT TACGICICAT TCCACAAGCC TCTAGTIGIA
CITIAAAAG CAAAGGITTA TAGICIGACA AIGCTAATTA TOCTAATTGI ATATAAAAA	360	15 ACCTICCICT CANTIGCTANT TCANTTAACA TACAGCCACG TAATCCAGAT GATATACAGT
THAMACATA GACCTITCTG THACAAAATT CTTAATCCTC TOGGTTGTAA TCATTACTTG	420	CAITCCAAAA TGATGTGTAA ACCITCGGGG TCCTTGAGGC CCAGCAGGAG CACTTCCTCC
CTACCAATIT ACATOCAACA TCTGCTAGGA CTGACATITG ATTITITITICC CCAAGAATGT	480	ATCAGGGTCA GCCGCGTTC CTTGGAGTCG CCCTTGTCGT CGTCGTCCTG CTCGTCGCGG
GTOAGTAGAT AAATGACATT TCAGAGCAGA TAITAATTTA CTIGTGGACA GAAAAAGAAA	540	COSCILETAGO COTOCITO GCTACTNACO GCGCCCCC CCGCCGCCC CTCCTTGTCG
CTCAAGATTO GTACTOGTCA CAAGCCTCTT CCCAATAGAA ATTATAAAAA CAGTAAGATA	009	GOGGOSTIGC GGAAGGCTC GGTGCGCCG
AAATTTAAAA AAAATCTAAA AAGGGGATGC ATAGGCAAAG AGTACCATAA ATGGCACAGC	099	25
TCAAAAAATC CCAGGACCAA TCAGACACAC ATCTTTTCTC TCTCCTTCAG CGACAAGAGG	720	(2) INFORMATION FOR SEO ID NO: 303:
TCGATTTTGC CATCAAATAA CCATGATTGA AGCAAGGGAG GGGCACCAGG TGTACAACTG	780	30 (1) SEDUENCE CHARACTERISTICS:
ATTAGATETT GENARATAET ANGATGGGAG CAGGGGTGGC CAGAAGAAGG GETAATTTAT	840	
ATATAATTCA AACTATATAC AGCATAAATG GAATGCAGCC CATCCCAAAC TGGCTCTGTG	006	000
aarcaattigg acctttatiag ttaaaattat aacaagigta ataatacaat agattiacat	096	35 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 303:
GOGNACCIAN ATCCANGGGA CATITITATAT TANGTATITA CTGTGCTGTT TCANTITANA	1020	GGGACGTOTG GTTTCAGCTC GTGGGCCTCC CCGTGGGTTT GCGACGTTTA GCGACTATTG
AATAATTTTG CTAAGTATAC ATCTCAACTG AAGTCTATGT AAAAAATGTC CTAATAGATA	1080	40 сасствосасе всясоваетв свявиствае всеятветте стветсова втантветна
CAGAMATTIA CCTITIGGIGA GITICAAGGCC TITTITGIGAC TICTIGICIGA ACTIGIAGGCA	1140	GOGETICCET GGGETICCAGG CTGTTGCGGG GTGTAGGTGG GAGTICACGA COGTTICGGG
GAATOCTAGA TOTACATOCA CATATGGAGA AACTCAAOCT GAGGTCATCC AAAAGCTGTG	1200	CCCGAGGTGT CCGCGAAGGT GGCGCACATG GGCGGCAGGG GAGAGCATGG CTCAGGGGAT
cotatgagga gactggaggt actttgaaag tcaaagtaga ccagaaaccc aaaacaggta	1260	45 GENCTIOGETIG GACCTIGAGA TGACAGGATT GGACATTGAG AAGGACCAGA TTATTGAGAT.
acacteagga togcaacagg gaatggaatg ccaatatogc actaaaactt tttttaaaa	1320	GOCCIGICTO ATARCIGACE CEGATOTICAA CATITITGGOT GAAGSTOCTA ACCIGALITAE
CAGAAAGAGG AAGGCCTCTC GTACCAGCAG AATCCTGTAC ACGTACAAAA AAGAAAAAGC		50 маласальска сателестве теследест стеленттее тотладелее атклеовель
caccaccat titgiaaac agaagccaat tatagtgigg gaaagtacaa attacagaaa	1440	STOTESCOTT ACCARGENG TRANSPARE TACANTACA TTGCAGCAGS CAGAGTATTGA
ACCAGAAGTC AACAGAAGAA AAACTACTGG TTTACTTGAG AGAAAGGAGA ATGGTTCACC	1500	ATTICTISTICS TITISTAGGAS ASCAGASCTICS TICCAGGGSCTIC TISTICCACTITG CAGGANATTIC
CCGAGCAGAG TIACTITGGTG AACGCCGCCA CCACCGCCCA CAGAACCTCA TIGGTGTTGG	1560	 AGTICATGAA GATAAGAAGT TTCTTGACAA ATACATGCCC CAGTTCATGA AACATCTTCA.
CCITCAGACA ITICACITICA GOSTICIAAGI CGAGAARINTO CCOCACTICIC ITIGGIACICA	1620	TTATAGATA ATTGATGTGA GCACTGTTAA AGAACTGTGC AGAGGCTGGT ATTCAGAAGA
AATCATACTG CTCGTCCAGA AGAGGGGAA AAGCATTCTC CAGGACGTCC GAGGCATGAG	1680	60 атапрааттт осассавара востретте театворска стратраса ттаргравая

 20 Ö 30 25 2 45 6 35 50 8 55 GCAGCCAATG CAGAAAATGG GTCAGCTCCT TIGAGAACCC CICCCCACCT ACCCCTICCT AGCTOTOGGE ACTITOTIGAAA GEAGAAGGEC AAGAACITOCT GGCCAGGACIT GCAAGGCTOT TOTTICOTAG ATGGGGTTTG CAGCIGCCAC TGAGCIGTAG CIGCGIAAGI ACCICCITGA AGTOCTOTT COCGCAGOTG CAGAGAGGAG GAAGACTATT AAAGGACAGT COTGATGACA ACAGCCAAAG CTGTTGCGGT GACCCTGCAG TCACACTGAC CCCACCTGAA AFTCTTGGCC CCCAGAGACA TCCCTGCCAT GCTCCCTGCT GCTCGGCTTC CCACCACCGT CCTCAACGCC TITAATICCC TTACCTACTC TIGCTCTATT TITTTATITG AAATGGAGAT GAGCAAAATA 2 ддададада лаадалалаа алааладссс ссисссс TOCTOTTTAT CTCTCCCACA TIGTCTTGCT AAATATAGAC TIGGTAATTA AAAAAAAAAA TTATATTTCT ATTCACATCA TTATTGAAAA TACCCAGCTC AGTGCCTGGC TTAATAAATG GGATGATCCT GAGCTGTTCA AACAAGCTGT ATATAAACAG ACAATGAAAC TCTTTGCAGA TOCATOCCAA AACCAATOCC TOCCAAACAA AATCTTAGAC ATCCCAATAT AATATOTTAG GAGGGGCTTG GTTTAAAAAA AGACCTTTCC CTCTCCCTGC CCCTAGAACA ACCAGTATTA CGAAGGACTC ATTCTTTCCT GAAGAAAGAG AAGAAAAATC GGACCTTCCT GAGACCACCG AAAGTAAAAA TGGAGCAACG TURGURARGY CURGATOGIC GTGTGGACAG CTGGCGARAC TTCCARGCCA ATACGRAGGG GGAAGAAGAG ATTGAAGCTC AAGAAAAAAGC CAAACGGGAA AGAGAGTGGC AGAAAAAACTT GCTGGAAATT AAAAGGAAAAG AGAGAGAAGC CAAAGAGATG CATGAAAGGA AACGACAAAG ACACATTCAT GGCTGAAGCA ATTITITIGGA CATTICTIGT TACCAAAAGA TCTATAATCA TEAGTEACCE CCCAAGGTCA CAGGCACAGA ACCTTTCCCC TOCATITION THICAATACG INFORMATION FOR SEQ ID NO: 305: E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (A) LENOTH: 1493 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AFTTAATATC GATCAGAGTA AFTCTTTTGT CCCACTTCCA CCCCAACATA GAGTAGTATT TGCTTTTTAG TOCTATORICC CITECTICITY ACATTGAAA1 1140 1500 1440 1380 1320 1260 1200 1537 360 300 240 180 120 420 600 540 480 8 780 720 660

20 25 15 5 330 45 6 S 55 50 CGARATACTA TITTICICCT ARTAIGCIGT TICCATIATG ACACAGCAGC ICCITIGIAA GAGGAAAATT ATAGAAAATG GGGAAAATGA GAAGACCGTG AGTTGATGCC AGTTATCATG GTACCAGGIC ATGICCATCC CTTGGTACAT ATATGCATTT GCTTTTAAAC CATTICTTTT CATCAAAGAG CITCAGITIT ACCGAAATAA CATCITCAAG AAAAAARTAG ATGAAAAGAA алдаалааа аалаалаала далаалаааа аалаалааа N CTTGGCAGAG CAGGCTTGGGA AAAGACAAGC CTCCAGGGCC TTCAGCTTGT ACGCCAACAT CGACATCCTC GAGGAAGAAG TAGACGCTGA TGCAGCTGAT GCAGCTGCTG CTGAAGAGGA GGATGGAGAG GREACATEGE AACTACAGEG GEGEEGGEGG GEGGEGEECEG AANAATEGAGE TGGCCCGGAA CCACATETTO CCCACTCCGC GCGCGGGGCT AGCGCGGGTT TCAGCGACGG GAGCCCTCAA CITITIONER RECOGCEGAR CECACETERE CICOACECTO GACGICTACE TICEGGAGGE 2 TICCTOGGCA TGAAGGGCTT TAAGGGACAG CTGAGCCGGC AGGTGGCAGA TCAGATGTGG ATGOATGATA CCTCAGGCTC TAGCTTCGAG GATATGGGTG AGCTGCATCA GCGCCTGCGC CTOGRETICA CICROTTOC TARCCTACTE CAROGARGA AGACGTETGA CACTATTARE CCTATICAAGA TOGTICAACTI CCCCCAGAAA ATTOCAGGTG AACTICTATOG ACCTCTICATO AGACCCTACT TIGATOTOGA GCCTGCTCAG GTGCGAACAG GGCTCCTGGA GTCCATGATC TOSOGAGGOT TOGAAGAAAA CATOCAGOGO GGAGGOTCAG CTOTGATTGA CATGGAGAAC TCATCCTTCA COGGAGGGCA CCCTGATGGG CACAGCCATT INFORMATION FOR SEQ ID NO: 304: (i) SEQUENCE CHARACTERISTICS: ž CHCTTCGGTT AACTTGCATC TCCAGATTGA TTACTCAAGC AGACAGCACA ARCGITATICE GOAGGCAACE TOEGGIGGET TETETETETICE ACGCIGATGG TITACTICCI IGCCIACCIG IGCAACGCCC AGAICACCAT GCIGCAGAIG SEQUENCE DESCRIPTION: SEQ ID NO: 304: (A) LENGTH: 1537 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double TOPOLOGY: linear GGCACCTGCT TCGGCTACTG GCTGGGAGTC 1140 1181 1080 1020 960 840 900 780 540 480 420 360 300 180 120 780 720 660 600

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ATCCACCICC ACCCCCTCTT CTACCICTTC TOGCTGTTGG TGGGTGGACT GTCCACACTG

900

TOGGCTATEG CCTCTTTEGG CATTECATTG TCCTCTTCAT CACCTATAAT

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COCAMBOTAG CADTOTTOOT GICTICGGACC GIGGGCCCCA CACAGCGGCT GCTCCTCTGT

960

GOCACCCTGG CTGCCCTACA CATGCTCTTC CTGCTCTATC TGCATTTTGC CTACCACAAA

GTONTAGAGG GGATCCTGGA CACACTGGAG GGCCCCAACA TCCCGCCCAT CCAGAGGGTC

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GARGETOCCA, CCATTGGTGC, TGCCTTCTCT, TCCCACAGCC, TGTAACTCAG, TGTTTTTGTAC	840		(2) INFORMATION FOR SEQ ID NO: 307:
TTCACTGAAT TOTGATGOTT AGAAACTTCG TGGATAGTTT GTGGAAATCA TCCAATTAAA	006	\$	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2860 base pairs
CATACTGCTT AAAACAGTGT TGCTGTGACT TCAGAGACAA GCCTGGAAGG GGCACCTTAG	096	•	(B) TYPE: nucleic acid (C) STRANDEDNESS: double
GAAGCCCCTT CGCTTCAGTT GCTGGCTTCT GGGTGTGCTC CCTTCGAAGG CCCAGATAAG	1020		
ACAGGSAACA CTTGTGAGCA CACAGAGCAG CATCTGATGC CCTGTGGTGT TTGGCATGTG	. 1080	10	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 307:
CCCCCTSTCT ACTGACGART CAGTSTGGCA TGAGGCCCAC GCCACCGAAA CCTTTCACTT	1140		GTOTINANCIG CTCTICNICAAT ATGGCTCCCC CGGGCTGGCA GRWRKTCRGT CMCKRGTCCC
TECAMAGAGE TRACESTECT CEACECAGTA CEATSTECTA GESTSTETE ATTISTIAGT	1200	51	TAGCCTOTOC TGACAGOGGA GAGTTAAGCT COCOTTOCC ACCOTOCOGG CTGGCCAGGT
GOTABTATIC TITIATOTATA ATAAATTITT ATACCCAAGC CATTGATGTA CITITICCTIG	1260	2	GOGCTGAGGG TGACCGAGAG ACCAGAACCT GCTTGCTGGA GCTTAGTGCT CAGAGCTGGG
TACTETICET TGTGGGTCC TTGTCTGGCT TGGCTGAACC CCAAATGCT TTGGGGTTGG	1320		GAGGGAGGTT COGCOGCICC TCTGCTGTCA GCGCCGGCAG CCCTCCCCG CTTCACTTCC
ACACACCTOS CTGAACCTTA GTITCTTCAT CTATGAANTG GGAATATGAA TTACTGCACC	1380	20	TECEGEAGCE CETIGETACTE AGAMGETECG GGATECEAGE AGECGECACG CECTIGGECTIC
AGCTITITAGS GCAGATITISC CATGGCATAT ACAAGSTAAC TACCATAGTS CTCCTTGGGT	1440		AGCTGCGGG GCTTCCAGTC AGGCCAACAC CGACGGGCAC TGGGGAGGAA GACAGGACCC
ATTGCCAATA TCCTATTATT TCTGTGTAAA ATGAAGATAC TGATTGTTTT GAG	1493	ν̈́	ITGACATOTO CATOTOGACA GAGGICCTIGG CTOGAAGCGA GCAGCOTOCT COTOCTAGGA
		3	TRACCICACC CICCAGCICT CCAGTITICA GGTIGGAGAC ATTAGATGGA GGCCAAGAAG
			ATGCCTCTGA GCCCAACAGA GCAAAGCTGG ATTTTGCCAG CGCCTCCCT CCCATGCAGT
(2) INFORMATION FOR SEQ ID NO: 306:		30	CACAGITICCA GGGCGAGGAC CGGAAATICG CCCCTICAGA TAAGAGICAA CCTCCAACTA
(1) SBQUBNCE CHARACTERISTICS: (A) LENGTH: 577 base pairs			COGNAGGG ACAGGTOCCA GTCAGCOGGA TOCANACCGA TITGACCGAG ATCGGCTCTT
(B) TYPE: nucleic acid (C) STRANDEDNESS: double		,	CANTGOGOTO TOCCOGANGA TCTGGCTGGA CTTCCAGAGT ACCTGAGCAA
		æ	GACCAGGAAG TACCTCACCG ACTTCGGAAA TACACAGAGG GCTCCACAGG TAAGACGGCC
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 306:			TGHIGAAGGC TGTGCTGAAA CCTTAAGGAC GGGGTCAATG CCTGCATTCT GCCACTGCTG
AATTICOGCAG AGGINATTATA TACACTATAC TGGCATTTAC TGTTTCACCC AGCCCGGAAA	09	40	CAGATICACE GGGACTETIGG CAATECTICAG COCCIDENIAA ATGOCCAGTIG CACAGATICÁC
CICAGAGATG TATATIGGAA AATTTACAAC TCCATCTACA TIGGITCCCA GGACGCTCTC	120		TATTACCGAG GCCACAGGG TCTGCACATC GCCATTGAGA AAGAGGAGTC TGCAGTGTGT
ATAGCACATT ACCCAAGAAT CTACAAGGAT GATAAGAACA CCTATATTOG TTATGAACTT	180	:	GAGCTECTG GTGGAGAATG GGGCCAATGT GCATGCCCGG GTCTGCCGGC ACTTCTTCCA
GACTATATOT TATAATITIA TTOTITATIT TOTOTITAAT GCACAGCTAC ITCACACCTT	240	45	GAAGGCCAA GCGACTIGCT TITATITICGG TGAGCTACCC CTCTCTITIGG CCGCTTCCAC
AAACTTOCTT TGAITTTGGTG ATGTAAACTT TTAAACATTG CAGATCAGTG TAGAACTGGT	300		CAAGCAGTGG GATGTGGTAA GCTACCTCCT GGAGAACCCA CACCAGCCCG CCAGCCTCCA
catagaggaa gaoctagaaa tocagtagca tgaittitaa ataacctgic tttottittig	360	50	GECCACTEAC TCCCAGGGCA ACACAGTCCT GCATGCCCTA GTGGATGATC TCGGACAACT
atottaaaca gtaaatgoca gtagtgacca agaacacast gattatatac actatactgg	420		CAGCTGAGAA CATTGCACTG GTGACCAGCA TGTATGATGG GCTCCTCCAA GCTKGGGSCC
AGGRITICA TITITAATIC ATCITIAIGA AGATITAGAA CICAITICCIT GIGITIAAAG	480	;	SCCYTCTGCC CTACCGTGCA GCTTGAGGAC ATCCGCAACC TGCAGGATCT CACGCCTCTG
ggaatottta attgagaat aaacatttgt gaacalaatg ytaaaaaaa aaaaaaaaa	540	55	AAGCTGGCCG CCAAGAAGG CAAGATCGAG ATTTTCAGGC ACATCCTGCA GCGGGAGTTT
araaaaaaa aaaaaaaaa aaaaaaaaa Aactiga	577		TCAGACTGA GCCACCTTTC CCGAAAGTTC ACCGAGTGGT GCTATGGGCC TGTCCGGGTG
		09	TOCOTOTATO ACCTOSCITIC TOTICACAGO TOTICAGAGA ACTCAGTCCT GAAGATCATT

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1380 1440

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539	

ATCCTOCTAG GGGGGATCTA CCTCCTCGTG GGGCCAGCTG TGGTACTTCT GGCGGGGCA CTGATCTACA TOTTCATCTT CACCOCTGTT GCCTACCATC AGCCTACCCT GAAGAAGCAG TOCAGCTOCT COAGTOCAAC TGATGGCCCA GATGCAGCAG GAGGCCAGAG GACAGAGCAG ATCTOGRAGO TGCAGAAAGC CATCTCTGTC CTGGAGATGG AGAATGGCTA TOGAGETETT CANATTENCE ATEGGENTOG GEGNGETOGE CTTCENGGNG CHGCTGCHET AGGAGGCTTG GCGCCCCGAA GCTCCTACAG GCCCCAATGC CACAGAGTCA GTGCAGCCCA GTICCAGCAC ACAGGCAGIC TACAGITICA IGWICCCIGA AGCCCIGGIG AGCCIGAGCC CONGOTIONS INCIGAGOS INSCINOSON SECTIONALES SCITTACIAN INCAASIOSA COCTOCTIVA CAGIOGIGIC CCAGGIOCIG IGTITICCIGG GCCANCGAGI GOIACCIGCO CONOTICATO TOGATOTOGT TOATAGACAG CTACTITICGA AATCOTOTIC CIGITOCAG GEOGEOCETE ACCIDADAGE GRAGGITOGA AACIOCATOC TOCTOACOGG CCACATOCTI TECTOSETTE CECTECEAAG GAGGATGAGG ATGGTGCCTC TGAGGAAAAC TATGTGCCCG TOCCTACOCT GIGIDAGGAC CCGICAGGGG CAGGIGICCC TCGAACICTC GAGAACCCTG CGATGAGCGC TOGTICCTTCA GGGTGGAGGA GGTGAACTGG GCTTCATGGG GAGCAGACGC TURACATOCT CATCOCCCTC ATGRAGOGRA COTCACAGTG TOGCCACTGA CAGCTGGAGO TECGEGGEAT GETGETGETG CTGCTGCTGG CETALGTGCT GETCACCTAC ATCETGCTGC ATATTITCAC TAACTCAAAA AAAAAAAAAA AAAAAAAAA AAAANGAGG GGGGCCCGKT AGGATETTIC CAACCACATE TGCTGGCTET GGGGTCCCAG TGAATTCTGG TGGCAAATAT AGGAAAAAGC AGCGGGCAGG TGTGATGCTG ACCGTTGGCA CTAAGCCCAG ATGGCAGCCC TOCIACOCIACA GCAGGACCIAG GGCAACGGGG CCCAGTACAG GGGTATCCTG GAAGCCTCCT ASCCAAWITC GCCCTATAAG TGAGTGCCWA TTACGATAAA (2) INFORMATION FOR SEQ ID NO: 308: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 308: SEQUENCE CHARACTERISTICS: (A) LENGTH: 876 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear TIGGIGGIGG 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2220 2160 2100 2040 1980 1920 1860 1800 1740 2860 8 35 30 25 20 2 5 45 50 55 8 COSTIGUCAC TOTOCTOCCO TOCCTCAGAG ACACCAAAACT GCCAAAAACA AGACGCGTAC CCCTGCCTGC CATACTGCGC CTAACTCGGT ATTAATCCAA AGCTTATTTT GTAAGAGTGA GETOCEGEE CEGGAETETO ATETETOTAG TOGECECETE CTECEGGGEE CETTITICGE TOCTOCTOGO ANCASOCCOO OCCCAMOTOC ATOGAGOTOT COTOCTAGGO GGCCTGCCCA CACCCOCAGE COCTCOCACA CCAGCGAGGG GGCCCACCTG GACATCACCC CCAACTCOGG CACCCGCAGC CGCTCCCCACA CCAGCGAGGG CACCCGAAGC CGCTCCCACA CCAGCGAGGG CATGACCCGC CTGATGCGAT CCCGCACAGC CTCTGGTTCC AGCGTCACTT CTCTGGATGG 2 GCTCTGOTGG AGACAAATGA GGTCTATTAC GTGGGTGCCC TCTCCAAAGG CGGGGTGGCG INFORMATION FOR SEQ ID NO: 309:

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SEQUENCE CHARACTERISTICS:

SEQUENCE DESCRIPTION: SEQ ID NO:

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480 420 360 (C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 2025 base pairs ઝ

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CTOCTOCAGG CGAAATOGGA TCTOCTCATC CCCAAGTTCT TCTTAAACTT CCTGTGTAAT OCCITICATI GUAAGAGCCC GUACCGACAC CGAATGGTCG TITTGGAGCC CCTGAACAAA

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CATEGRACIS CIOCISCISC IGGCCIACGE SCIOCICACE TACATOCISC IGCICAACAT TIBOCOCCCC GAAGCICCIA CAGOCOCAA TOCCACAGAG TCAGTGCAGC CCATGGAGGG CAGCACACAG GCATCTACAG TOTCATGATC CAGAAGCCCT GGTGAGCCTG AGCCAGGANN CIGCITOTOT CICCOCIGGI GCIGGGCIGG CIGAACCIGC TITACIATAC ACGIGGCITC CTTCCCCTCC CAAGGAGGAT GAGGATOGTG CCTCTGAGGA AAACTATGTG CCCGTCCAGC AGCGCTOGTG CTTCAGGGTG GAGGAGGTGA ACTGGGCTTC ATGGGAGCAG ACGCTGCCTA GGAAGCTGCA GAAAGCCATC TCTGTCCTGG AGATGGAGAA TGGCTATTGG TGGTGCAGGA OCTICATICOCIC CTICATIGNAGIC GAGACICOMICA ACAGTOTICOC CACTIGACIAGIC TOGAGICATICT CTICAAATIC ACCATCGGCA TOOGCGAGCI GGCCTTCCAG GAGCAGCTGC ACTTCCCCGG ACAGGAGGAC GAGGGCAACG GGGCCCAGTA CAGGGGTATC CTGGAAGCCT CCTTGGAGCT ТСАСТААММ ААЛААЛААА АААЛААЛАА АСТОБА TOCTOCAGIO CAACIGAIGG COCAGAIGCA GCAGGAGGCO AGAGGACAGA GCAGAGGATO AGAAGCAGCO GGCAGGTGTG ATGCTGACCG TTGGCACTAA GCCAGATGGC AGCCCCGATG TITICOAACCA CAICIGCIGG CICIGGGGIC CCAGIGAATI CIGGIGGCAA ATATATATITI GRACCEGTICA GOOGLAGGTG TECETICGAAC TETEGRAGAAC CETGTEETIGG 840 660 600 540 480 420 360 300 240 876 780 720 180 120 69

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~	CTGGTCAGAA GOCCAGCCCC CCACTTCCCG ITTCCTCTTT AACTGAGGAG AAGCTGATCC	099	
,	AGITICCGGA AACAAAAICC TITICICATI TGGGGAGGGG GGIAATAGIG ACAIGCAGGC	720	
	ACCTICTTTTA AACAGGCAAA ACAGGAAGGG GGAAAAGGTG GGATTICATGT CGAGGCTAGA	780	
10	GOCATTIGGA ACAACAAAIC TACGIAGTIA ACTIGAAGAA ACCGAITITIT AAAGTIGGIG	840	
	CATCTAGAAA GCTTTGAATG CAGAAGCAAA CAAGCTTGAT TTTTCTAGCA TCCTCTTAAT	006	
7	GTGCAGCAAA AGCAGGCRAC AAAATCTCCT GGCTTTACAG ACAAAAATAT TTCAGCAAAC	096	
2	OTHOGOGATIC ATGGTTTTTG AAGGCTTTAG TICTIGCTTTC TGCCTCTCCT CCACAGCCC	1020	
	AACCTCCCAC CCCTGATACA TGAGCCAGTG ATTATTCTTG TTCAGGGAGA AGATCATTTA	1080	
20	GATTICITIT GCATTCCTTA GAATGGAGG CAACATTCCA CAGCTGCCCT GGCTGTGATG	1140	
	AGTOTECTTO CAGGGCCGO AGTAGGAGCA CTGGGGTGGG GGCGGAATTO GGGTTACTCG	1200	
25	ATOTIAAGGGA TICCTIGTIG TIGTOTIGAG ATCCAGTGCA GITGTGATIT CIGTGGATCC	1260	
}	CAGCTIGGT CCAGGAATT TGIGTGATTG GCTIAAATCC AGTTTTCAAT CTTCGACAGC	1320	
	TOOCTIOGNA COTGNACTCA GTACCTGNAC CTGTCTGACC COOTCACOTT CTTGGATCCT	1380	
30	CAGAACTETT TOCTETTOTE GOOGTOOGG TOGGAACTEA COTGOGGAC GOTGOCTGAG	1440	
	AAAAIGIAAG GAFTCTGGAA TACATAFTCC ATGGGACTFT CCTTCCCTCT CCTGCTTCCT	1500	
35	CTITICCISC TCCCTAACCT TICGCCGAAT GGGGCAGCAC CACTGACGTT TCTGGGCGGC	1560	
3	CAGIGGGGCT GCCAGGITCC TGTACTACTG CCTTGTACTT ITCAITITIGG CTCACCGTGG	1620	
	ATTITICICAT AGGAAGITTO GICAGAGIGA ATIGAAIRIT GIRAGICAGC CACIGOGACC	1680	
40	CGAGGATTIC TOGGACCCCG CAGTTGGGAG GAGGAAGTAG TCCAGCCTTC CAGGTGGCGT	1740	
	GAGAGGCAAT GACTCGTTAC CTGCCGCCCA TCACCTTGGA GGCCTTCCCT GGCCTTGAGT	1800	
45	AGAAAAGTOG GGGATCGGGG CAAGAGAGGC TGAGTAGGGA TGGGAAACTA TTGTGCACAA	1860	
?	GICTITICCAG AGGAGTITCT TAATGAGATA TITIGIATITA TITICCAGACC AATAAATITIG	1920	
	TAACTTTGCA AAAAAAAA AAAAAAAA AAAAAAAA AAAAAAAA AAAA	1980	
20	GAGGGGGGC CGTACCCAAT TCGCCGTATA TGATCGTAAA CAATC	2025	

(2) INPORMATION FOR SEQ ID NO: 310:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3026 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 310:	
2	TAGGAGGAC TGAAATATCC TAACCCCCTA AGCTCCAGGT GCCCTGTGGA ACGAGCAACT	9
	GOACTATAGC AGGOCTGGGC TCTGTCTTCC TGGTCATAGG CTCACTCTTT CCCCCAAATC	120
2	TTCCTCTGGA GCTTTGCAGC CAAGGTGCTA AAAGGAATAG GTAGGAGACC TCTTCTATCT	180
2	AATCCTTAAA AGCATAATGT TGAACATTCA TTCAACAGCT GATGCCCTAT AACCCCTGCC	240
	TOCATTICT CCTAITAGGC TATAAGAAGT AGCAAGATCT TTACATAATT CAGAGTGGTT	300
15	TEATTGCCTT CCTACCCTCT CTAATGGCCC CTCCATTTAT TTGACTAAAG CATCACAG	360
	TOSCACTAGE ATTATACEAA GAGTATGAGA AATACAGTGE TTTATGGCTE TAACATTACT	420
ç	OCCITICAGIA TCAAGGCTGC CTGGAGAAAG GATGGCAAGCC TCAGGGCTTC CTTATGTCCT	480
3	CCACCACAAG AGCTCCTTGA TGAAGGTCAT CTTTTTCCCC TATCCTGTTC TTCCCCTCCC	540
	COCTCCTAAT GSTACGTGGG TACCCAGGCT GSTTCTTGGG CTAGGTAGTG GGGACCAAGT	009
22	TCATTACCTC CCTATCAGTT CTAGCATAGT AAACTACGGT ACCAGTGTTA GTGGGAAGAG	099
	CIGGOTITIC CINGININC CACIGCATCC TACTOCTACC TGGTCAACCC GCTGCTTCCA	720
Ş	GETATIGGGAC CTOCTAAGTG TGGATTACC TGATAAGGGA GAGGGAAATA CAAGGAGGGC	780
3	CTCTOSTIOTT CCTGGCCTCA GCCAGCTGCC CACAAGCCAT AACCAATAA AACAAGAATA	840
	CTEAGLEAGT TITITATICTG GGITCICTIC ATTCCCACTG CACTTGGTGC TGCTTTGGCT	900
35	GACTOGGIAC ACCCATAAC TACAGAGTCT GACAGGAAGA CTGGAGACTG TCCACTTCTA	096
	GCTCGGAACT TACTGTGTAA ATAAACTTTC AGAACTGCTA CCATGAAGTG AAAATGCCAC	1020
5	ATTITIGETTT ATRAITTICTA CCCATGITIGG GARARACTIGG CTTTTTCCCA GCCCTTTCCA	1080
}	GOSCHIAMAA CICAACCCCT TCGAIAGCAA GICCCAICAG CCIAITAITI ITTIAAAGAA	1140
	AACTIGCACT TGTTTTTTT TTTACAGTTA CTTCCTTCCT GCCCCAAAAT TATAAACTCT	1200
45	ANGTOTANAN ANANGTCTTA ACAACAGCTT CTTGCTTGTA AAAATATGTA TTATACATCT	1260
	GINITITIMA ATTICIOCICC TGANAANIGA CTGTCCCATT CTCCACTCAC TGCATTITGGG	1320
Ş	GCCTTTCCCA TTGGTCTGCA TGTCTTTAT CATTGCAGGC CAGTGGACAG AGGGAGAAGG	1380
3	GAGAACAGGG GTCGCCAACA CTTGTGTTGC TTTCTGACTG ATCCTGAACA AGAAAGAGTA	1440
	ACACTGAGGC GCTCGCTCCC ATGCACAACT CTCCAAAACA CTTATCCTCC TGCAAGAGTG	1500
25	GOCTITICCAG GGICTITACT GGGAAGCAGT TAAGCCCCCT CCTCACCCCT TCCTTTTTTC	1560
	TITICITIACT CCITIGGCIT CAAAGAITT TGGAAAAGAA ACAATAIGCT TIACACICAT	1620
9	TITICARITIC TARATTIGCA GOGGRIACTO ARABATAGOG CAGGIGGCCT AROCTICGIG	1680

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6 ઝ છ 25 20 45 2 5 Ś TAATGICCIT CCATCTOTGA GOTGAYAGGC AAGGATGAAA GACAAAGAAG GAAAAAGAOTA TCAAAAGGCAG CCTTAAACTG ACGCTACAMA ATAAACCTGG GCAAGTGAGG CAAGAGAAAT GAGGAAGAAT CTGGAGACAG TAACATTICA TIAACCAAAG AAAGIGGGIC ACCIGACCIC IGAAGAGCIG TITOGCCITOTO AGAGCTIGAT TAGAAGCCAA GACAGIOGCA GCAAAGGAAG ACTITOGCCCA CACCCAAGGT CANCCAAACA ACTIGGITGT GAACCCAACT GCCTTAACCT ICTGGGGGAG AGATCATTOT ACAAGTAATT TIGCACAGAC AICICCICAC CCCAGIGCCI GICIGGAGCI ATTTATAAAT TIGAAATCCA AACTACTTIC TIAATATCAC TIIGGICICC ATTITICCCA AAAAGATTCA AAGCTCTAAT AGAGTCACAG CTTTCCCAGG TATAAAACCT AAAATTAAGA AATTCTAAAA GAGAAGGGAG CTGAGGCCAT TCCCTGTAGG AGTAAAGATA AAAGGATAGG AGTACCTOTA AGCATTITAG GTCCCAGAAT GGAAAAAAAA ATCAGCTATT GGTAATATAA AAAGGAGATC ATTTAGITGG GTCTGAAAGG AAAAGICTTT GCTAICCGAC ATGTACTGCT AGTACTCAGG CCACTCCAAT CACCCTACAA GATGCCAAGG ARAGRACART AGRACTOGIC TICCRITITIG CCACCITICC IGITCRIGRC ROCTRCIRAC TAAAGTTGAG GGGAGAGGAA ATCTTAAGAT TACAAGATAA AAAACGAATC CCCTAAACAA AATTINCCAA ATAGAGATNG TATTAC ATGTAAAACA GAATATTCTG TAAACCTAAT GTCTGTATAA ATAATGAGCG TTAACACAGT GGGTGCATGG AGGAATTGGG ACCTGGTTAT GTTGTTATTC TCGGACTGTG AATTTTGGTG GGAAAAACCT GIGGGTIGIG CTAATTICIG ICCAGAAAAT AGGGIGGACA GAAGCTIGIG GGACAGGAAA TATGTCCCCC CCTAACTTTC TTGCTTCAAA AATTAAAATC CAGCATCCCA AGTACAATAA GCAGAGGTGG AAAATGATCT AGTTCCTGAT AGCTACCCAC AGAGCAAGTG AAAATATTCA ATAAGAAGTC AAAAAAAAAA AAAAAAAACT CGAGGGGGGG CCCGGTACCC GGGGATTAGC TAGACTAGGA GACCCAGAAG TGAATGGGAA AGGGTGAGGA CTTCACAATG TOCOTOGRAGI CAGITITITITI AAAAAGITAA CICITAGITI TIACITGITI AGGTCCCAGG AAGTCCAGCT 2880 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2160 2100 2040 1980 1920 1860 1800 3026 3000 2940 2220

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INFORMATION FOR SEQ ID NO: 312:

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SEQUENCE CHARACTERISTICS:

20

TCAATAAAGC CACATGTGCC TGTGGCCCAA AAAAAAAAA AAAAAAAAA AAAAAAAAA

AACTOGAGOG GOGGCCCGGT ACCCAAATCG CCGGATATGA TCGTAAACAA TC

712 660 CHOSCIGATE GOTTOGGGCC ACATECETEC TECTTECTEC TOCTCATECT CTCTGCCTTT

540 480 420 360 300 240

600

CONSTITUTO ACCIGGACOT AGCACOCAGO ACCITITOTOT CAGOIGAGIG GOIGGAGIGG

15

ACAGCCTTCT

5

GOSTICAGOOT TOCTIGAGOOT ATIGICTIGGAG CACTITOTIGG GAGGOOTIGGT CACCADAGTO

ACCTICACIG GGAIGAIGCG CIGCAGCCAG CIGGCCCCCA GGGCCIGCAG GCCACACACI

GGCCACGCTG GAGCTGCTGG GGAAGCTGCT GCTGGGCACT CTGCGGAGGC

GCCTTGOTCT TCCACCTGGA CACCCTGGGG GCCAGCATGG ACGCTGGCAC AATCTTGAGA

CIBCIBCCIC IGITEARGIC GGIOCIBCGC TICCGCCTCG GGGGCCIAGC CIGICAGACT GIBGICIBET CCATEBEIGG CICCICCEIG GOIGGGACET IGEIGGCCAA GCACIBGAAA CICCIBCIDG ACCACBBOT TICIBCICCO GASTIBBBAC IGIBBAAIGG IGIBBBIBCI

> 180 120

45 6 35 8 SS 8 THAGACTIAT TAATAIGTIC TIGTCCIGTA TITATACATA TGIGTATTIT GGAAAGUATT GITATIGGGT TITIGGTIGG TITITIGITIG TITITITACIA IGCTITGGIC IGTAAAAAIA TIGCCAACCT ICIGGITGAG CIGCAAGAAA ATATTTATGG IGAGAACTIT CAAAATTICA GAACTIICAG GAGGGCAAGA GAATATCAAA CAAAGATIIC TGGAAGTATT TOTOTTOTTT TOCCTOGATG AMARTATCAG TATTAAGTAG ACAGCATATT ATTCAAGTGT ACATAATTCT GATGAGGAAA AAAAATCTTT GCAATTCTTT AAGCCATATT GTTGTTTTTC TOCAACTGAA CTACATTCAG AAGGAAATAT TOTCTACATA GAATATTATA TGAAGTTGGT GCCTTTTTTA AGOGAAGCTA TAATICGATA CATAGIGAAA AAGGGAATGG TGACCCCTTT GAGGGCTCTC AATCAGCAGG GCCCCAGGAG GGAAGAAGAA GTGGGGCAAA GCCTGGCCTC TCTTATCTTG TTATTACGGT TTTATTAATT TTGTAGAGGG ACAGGGAGTG GGCAAGGGG GIOCCICITO CACIGAGGAT AACAAACAGC ATIGIAATCC ATICICITOC ACCITCITCI AGAAGCAGCT TATTTGACTA ACCAGCCCCT CTGTGGTCCA CCAGCGTCTT GGCTTGGTGG £ SEQUENCE DESCRIPTION: SEQ ID NO: 312: (A) LENGTH: 1289 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear Terentice

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720 660 600 540 480 300 240 180 120 60

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) TYPE: nucleic acid
) STRANDEDNESS: double
) TOPOLOGY: linear LENGTH: 712 base pairs 50

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 311:

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GCAGGCTTTG TOCTCACCTA CAAGCTGGGT GAGCAGGGTG CCAGCAGCCT GTTTCCTCTT

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 311:

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1289	CTTGAGGGGG GENCCGGTAC CCAATTNTC	C
1260	aatatagcct cttcatgctg tattaaaags acttttaaaa gcaaaaaaa aaaaaaaa	
1200	GOGATIGGCAT TCCCTTTOTGT GCCTTATTCC TGGAGAATCT GTATACGGCT CGCCTATAGA	8
1140	COTOCOCCA GCTCGGAAGG TGCGTGGAGA AGGCTCCGAC GTCTCCGAAG TGCAGCCCTT	
1080	CCCGTCCGTG TCTGTGTCTG TCCATAGGTG TGAGTCCAGC TAAAAAGACA AAACAGAACC	•
1020	GGCTTATICCT TCTCTGTTGC CAACCTTGCC GTCCGACCTC CTCCGCCCCC ATGCGGTGAC	C
960	ATGGGGGTGT TOCAGOCTTC AGGGAGATGG CCAAGGGGTC COCTGGGGGC TGTGGCAGGG	
900	TAAAGGAATA TITIAATCCAA CCTCACTACA TIGTAGCTCA GTCCAAGGAC TAAGCCTGAA	3
840	GANATOTICAA COGGACCOOT TACACCAGOC CTCCAGCATO TAATAGACTT GAATOTACTO	
780	OCCOCTOGGO AGCTITIGOCA TOTGAGCCAC OCCTCCTCCA GGCCATGCTC CTTGAACTTG	

(2) INFORMATION POR SEQ ID NO: 313:

(i) SEQUENCE CHARACTERISTICS:
(i) SEQUENCE CHARACTERISTICS:
(i) LENGTH: 22 amino acids
(ii) TYPE: amino acid
(iv) TYPE: amino acid
(iv) TYPE: amino acid
(iv) TYPE: amino acid
(iv) SEQUENCE SESCHITCH: SEQ ID NO: 313:

Met Phe Leu Ile Phe Val Tyr Phe Leu Lys Ile Leu Phe Ser Ser Ser 1 1 5

35 Leu Pro Phe Leu Trp Leu 20 40 (2) INFORMATION FOR SEQ ID NO: 314:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 128 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(xi) Sequence description: Seq id No: 314:

Met Het Phe Leu Thr Gin Gly Gly Pro Leu Pro Ser Thr Arg Ala Arg

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Pro Thr Cys Gin Ala Gly Ala Leu Pro Lys Pro Ser Gly Leu Leu Gly
20 20 30

Val Thr Cys Trp Asn Gly Leu Lys Gly Pro Leu Cys Gly Asn Arg Cys
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46
48
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Ser Pro Asn Thr Leu Leu Leu Ala Ala Arg Gln Ala Leu Trp Lys Gly
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60 Arg Gly Arg Thr His Gln Asp Leu Pro Gly Pro Leu Gln Gly Arg Gln

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65 70 75 80 Even Gly Pro Glu Pro Lys His Leu Ala Leu Leu Pro Pro Arg Gly Gln 85 85 90 95

Pro His Ile Asn Cys Thr Val Phe Ser Leu Lys Ala Ser Phe Ile Lys 125

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Glu Ala Ser Trp Ala Ser Ser Leu Pro Gly Gln Gly Pro Leu Pro Leu 100

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) INFORMATION FOR SEQ ID NO: 315:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 28 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 315:

25 Met Gin Phe Leu Leu Thr Ala Phe Leu Leu Val Pro Leu Leu Ala Leu 1 5

Cys Asp Val Pro Ile Ser Leu Gly Phe Ser Pro Ser 20

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(2) INFORMATION FOR SEQ ID NO: 316:

(i) SEQUENCE CHARACTERISTICS:

(A) LEWITH: 64 antho acida

(B) TYPE: anino acida

(D) TOPOLLOSY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 316:

Met Asp Gly Phe Ser Ser Arg Leu Phe Ser Ser Leu Pro Phe Val Ala 1 5

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Ala Cys Cys Tyr Gln Thr His Cys Ser Leu Xaa Gln Leu Ser Ser Ala 35 45 Phe Ser Xaa Wet Gly Glu Ser Cys Val Gly Glu Arg Glu Tyr Xaa Phe

50 Phe Ser Xaa Met Gly Glu Ser Cys Val Gly Glu Arg Glu fyr Xaa Phe5050

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(2) INFORMATION FOR SEQ ID NO: 317:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 21 amino acids

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Lys Gln Asp Lys Lys 20 Met Pro Leu Ile Asn Leu Leu Leu Leu Tyr Tyr Val Pro Asn Gly Gly
1 10 15 Ser Gly Thr Pro Val Pro Glu Asn Val Ile Cys Gly Val Thr Lys Gly $20 \\ 20$ Met Leu Trp Trp Ser Arg Asp Tyr Thr Met Val Phe Leu Leu Phe Thr 1 15 Pro Gln Gly Lys Lys Lys Lys 35 Met Gly Arg His Leu Val Leu Val Met Phe Ile Thr Thr Ser Leu His 1 10 15 (2) INFORMATION FOR SEQ ID NO: 318: Met Val Phe Thr Gly Asp Leu Val Ile Arg Gly Arg Thr Glu Leu Ser $20 \ \ 25$ Met Val Cys Ser Ser Leu Cys Asp Ile Gly Gly Ile Ile Thr Pro Phe (2) INFORMATION FOR SEQ ID NO: 320: INFORMATION FOR SEQ ID NO: 319: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 318: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 317: (i) SEQUENCE CHARACTERISTICS: (1) SEQUENCE CHARACTERISTICS: (i) .SEQUENCE CHARACTERISTICS: (X Ĕ. SEQUENCE DESCRIPTION: SEQ ID NO: 319: (B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 320: (A) LENGTH: 33 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 39 amino acids (A) LENGTH: 88 amino acids 33 30 25 5 3 8 5 8 S 8

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The Val Phe Arg Leu Arg Glu Val Trp Gln Ala Leu Pro Leu Ile Leu 20 25 30 Gln Thr Ser Glu Pro Ser Gly Thr Glu Thr Lys Gly Val Ala Leu Pro Glu Thr Met Lys Asp Ala Glu Asn 50 Phe Ala Vai Leu Gly Leu Leu Ala Ala Gly Val Thr Leu Leu Leu Pro 35Leu Gly Arg Lys Ala Lys Pro Lys Glu Asn Thr Ile Tyr Leu Lys Val 65 70 75 80 10 5

8 (2) INFORMATION FOR SEQ ID NO: 321:

Ē (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 321: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENOTH: 23 amino acids

Met Gln Pro Gly Ala Gly Val Leu Val Leu Gly Leu Leu Leu Pro Pro 1 15

Pro Gln Ser Pro Ser Leu Ser 20

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 27 amino acids

(2) INFORMATION FOR SEQ ID NO: 322:

Met Thr Phe Thr Leu Gly Asp Ser Gln Val Leu Leu Ile Asn Leu Phe
1 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 322:

(D) TOPOLOGY: linear

Pro Ser Met Pro Ser Gly Ser Cys Ala Arg Pro 20 25

(2) INFORMATION FOR SEQ ID NO: 323:

Ě (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 323: (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 64 amino acids

Met Cys Leu Glu Cys Trp Ala Glu Asn Leu Gly Pro His His Thr Ser

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Pro Ser Ile Pro Ala Met Phe Pro Asn Lys Ser Leu Leu His Cys Ile Phe Xaa 12 Ser Ser Gly Cys Phe Gln Gln Gln Glu Met 35 40 ន Leu His Phe Val 55 Ser Leu Leu Asn Pro Arg His Leu Cys Leu Phe Val

INFORMATION FOR SEQ ID NO: 324:

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Ser 50

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Val

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(D) TOPOLOGY: linear (xi) ನ

SEQUENCE DESCRIPTION: SEQ ID NO: 324

Leu Ser Thr Ser Glu Tyr Ser Gln Ser Pro Lys Met Glu Ser Leu 5 Ser Ser His Arg Ile Asp Glu Asp Gly Glu Asn Thr Gln Ile Glu Asp 20 Het 22

Ser Ile Leu Met Asn Pro Ala Gln Asp Gly Glu Val Gln Leu 55 Pro Met Ser Pro Val Leu Asn Ser Lys Phe Val 35 40 45 Asn Asp S 35

Pro Ala Glu

Thr Glu

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Lys Thr Lys Gly Asp Asp Thr Asp Thr Arg Asp 70 75 80 Asp Ile Ser Ile Leu Ala Thr Gly Cys Lys Gly Arg Glu Glu Thr Val 90 95 Ser 65 各

Gln Asn Asp Asp

Ala Glu Glu Val Cys Ile Asp Leu Thr Cys Asp Ser Gly Ser Gln Ala 100 Val Pro Ser Pro Ala Thr Arg Ser Glu Ala Leu Ser Ser Val Leu Asp 115

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Glu Glu Ala Met Glu Ile Lys Glu His His Pro Glu Glu Gly Ser 130 20

Glu Glu Leu Lys Glu Glu Asn Met Glu Ser Val Pro Leu His Leu 175 Gly Ser Glu Val Glu Glu Ile Pro Glu Thr Pro Cys Glu Ser Gln 150 150 55

Ser Leu Thr Glu Thr Gln Ser Gln Gly Leu Cys Leu Arg Arg His Pro 180 8

Ŋ Lys 195 Lys Lys

(2) INFORMATION FOR SEQ ID NO: 325:

(D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 325: (A) LENGTH: 252 amino acids (1) SEQUENCE CHARACTERISTICS: TYPE: amino acid

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Met Gly Gly Asp Leu Val Leu Gly Leu Gly Ala Leu Arg Arg Lys Arg Leu Leu Glu Glu Lys Ser Leu Ala Gly Trp Ala Leu Val Leu 20 2

Ala Xaa Xaa Gly Ile Gly Leu Met Val Leu His Ala Glu Met Leu Trp Leu Ser Asp Phe Gly Gly Cys Ser Ala Val Asn Ala Thr Gly His 2

Leu Trp Leu lle Pro lle Thr Phe Leu Thr lle Gly Tyr Gly Asp Val 65

22

Va] Met Gly Val Cys Cys Thr Ala Leu Leu Val Ala Val Val Ala Arg Lys 100 5,50 겵 Š ile Val Cys Leu . 90 ĘŽ Trp Gly Met Val Pro Gly Thr

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ABP Leu Glu Phe Asn Lys Ala Glu Lys His Val His Asn Phe Met Met 115 35

Ile Gln Tyr Thr Lys Glu Met Lys Glu Ser Ala Ala Arg Val Leu Gln 130 Glu Ala Trp Met Phe Tyr Lys His Thr Arg Arg Lys Glu Ser His Ala 145 130 6

Ala Arg Xaa His Gln Arg Xaa Leu Leu Ala Ala Ile Asn Ala Phe Arg 175 Gln Val Arg Leu Lys His Arg Lys Leu Arg Glu Gln Val Asn Ser Met 180

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Val Asp Ile Ser Lys Met His Met Ile Leu Tyr Asp Leu Gln Gln Asn 205 Ser Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp Thr Leu Ala 210 Ę

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Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly Pro 225 235

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Ser Lys Xae 250 Arg Gln Leu Pro Glu Pro Ser Gln Gln 245

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Asn Lys Leu Xaa 65 Glu Lys Thr Thr Glu Asn Lys Glu Ser Asn Pro Phe Ile Leu Gln Val 50 55 lle Val Ser Cys Arg Lys Asp Gly Pro Asp Ala Ala Ala Ala Pro Ala 20 Met Trp Arg Cys Arg Gly Lys Leu Ser Phe Pro Leu Phe Ala Val Val 1 5 Thr Val His Thr Cys His His Gln Ala Phe Leu Val Leu Ile Gly Trp \$45\$Ala Cys Trp Glu Gly Val His Ser Glu Pro Val Cys Arg Thr Val His 20Met Gly Glu Gly Lys Asn Gly Phe Gly Gly Phe Val His Thr Ala Asp $1 \ \ \, 1$ Val Ile Lys Asn Asn Ser His Tyr Gln Thr Ser Lys Ala Leu Glu Leu 35 Ser Lys Ser Gly Lys Glu Arg Lys Glu Ala Phe Leu Thr Ala Ile Ile 50 55 (2) INFORMATION FOR SEQ ID NO: 327: Lou Asn Ser Arg Ser Ile His Ile Ser Cys Ser Trp Pro Pro Ser Pro 65 70 75 80 Val Pro Gln Xaa (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 326: 9 ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 327: (A) LENGTH: 84 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear

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Phe His Arg 35

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35

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INFORMATION FOR SEQ ID NO: 328:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 36 amino acids

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(2) INFORMATION FOR SEQ ID NO: 326: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 68 amino acids

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(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 amino acids

(2) INFORMATION FOR SEQ ID NO: 331:

552

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Met Leu Leu Ile Asn Leu Leu Trp Leu Val Thr Met Ile Lys Ser Val 1 15 Ile Asn Asn Asn Ile Ile Leu Phe Leu Lys Lys Lys Ser Leu Phe Phe $20 \ 20 \ 30$ Leu Glu Ser Tyr Asn Trp Lys Val Ser Cys Gln Leu Arg Glu Xaa 50 S5 60 Asp Met Thr Val Ile Leu Arg Gly Arg Ala Gln His Lys Thr Ala Met \$35\$Leu Ile Pro Gly Glu Ser Arg Leu Ala Pro Thr Phe Asn Pro Ser Ala
20 25 30 Met Thr Phe Pro Phe Glu Lys Lys Ile Val Ala Phe Ser Ala Phe Tyr 1 5 10 Met His Ser Lys Gly Ser Ser Leu Leu Leu Phe Leu Pro Gln Leu Ile 1 10 15 Leu Ile Leu Pro Val Cys Ala His Leu His Glu Glu Leu Asn Cys Cys 25 30 (2) INFORMATION FOR SEQ ID NO: 330: INFORMATION FOR SEQ ID NO: 329: (B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 328: (i) SEQUENCE CHARACTERISTICS: (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 329: SEQUENCE DESCRIPTION: SEQ ID NO: 330: (A) LENGTH: 63 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 35 amino acids

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Ile Asp Ser Val 35

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Met Gly Ala Leu Val Leu Leu Leu Cys Leu Leu Val Gly Val Gln Gln (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 331: (B) TYPE: amino acid (D) TOPOLOGY: linear Ser Gly Ser Val Trp Asp Ser 20 S

(2) INFORMATION FOR SEQ ID NO: 332:

9

(A) LENGTH: 40 amino acids SEQUENCE CHARACTERISTICS: TYPE: amino acid (D) TOPOLOGY: linear 2

SEQUENCE DESCRIPTION: SEQ ID NO: 332: ž

Met Gin Ser Ala Glu Ile Leu Ser Trp Thr Asp Val Leu His Asp Phe 1 5 10 20

Leu Phe Leu Trp Pro Ala Phe Glu Asp Arg Ala Leu Leu 20 30 Leu Phe Ser 25

Ile Phe Thr Leu Asn Gln Ile Val 35

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(2) INFORMATION FOR SEQ ID NO: 333:

(A) LENGTH: 111 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 35

Met Gln Ser Leu Val Gln Trp Gly Leu Asp Ser Tyr Asp Tyr Leu Gln 10 15 15 SEQUENCE DESCRIPTION: SEQ ID NO: 333: (ž 6

Asn Ala Pro Pro Gly Phe Phe Pro Arg Leu Gly Val Ile Gly Phe Ala 20 45

Gly Leu Ile Gly Leu Leu Leu Ala Arg Gly Ser Lys Ile Lys Lys Leu 35

Gln Gln Ala Ile Val Phe Ala Gln Val Ser Gly Glu Arg Leu Tyr Asp 65 79 Val Tyr Pro Pro Gly Phe Met Gly Leu Ala Ala Ser Leu Tyr Tyr Pro 50 20

Trp Gly Leu Arg Gly Tyr Ile Val Ile Glu Asp Leu Trp Lys Glu Asn 95 55

Phe Gln Lys Pro Gly Asn Val Lys Asn Ser Pro Gly Thr Lys Xaa 110 8

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(2) INFORMATION FOR SEQ ID NO: 334:

(A) LENGTH: 106 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS

SEQUENCE DESCRIPTION: SEQ ID NO: 334: Œ.

2

Met Ala Pro Ser Leu Leu Leu Ala Pro Leu Cys Ser Leu Glu Ala

Ser Pro Leu Glu Lys Gln Cys Gln Leu Pro Gly Ile Phe 20 30 Ser Ala Gln Leu Leu ž Pro Leu Leu L 40 Cys Gln Leu Gln Leu Pro Cys Val Leu Ser

2

Lys Gly Ile Val Xaa Pro Arg Cys Pro Ala Ser Leu Pro Gln Pro Pro 50

2

Leu Pro Leu His Cys Thr Glu Arg Xaa 75 Trp His 70 Pro Ala Pro Ser His 65 25

Pro His His Leu Pro Leu Gln Gly Gly Ser Ser Asn Met Glu Glu Xaa 85

Asn Tyr Arg Gly Tyr Xaa Asp Ala Gln Leu 100

2

(2) INFORMATION FOR SEQ ID NO: 335:

35

(A) LENGTH: 50 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: 3

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 335:

5

Met Thr Thr Cys Leu Phe Gly Leu Leu Ser Cys Glu Met Ser Ala Gln 1 5

Val Ser Gln Lys Ser Cys Val Tyr Asp Glu Ser Glu Cys Phe Ser Ser 20 45

Val Gly Gln Leu Leu Ala Leu Leu Ile Leu Val Tyr Val Leu Pro Ser 15

Ile Xaa S0

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(2) INFORMATION FOR SEQ ID NO: 336:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 48 amino acids (B) TYPE: amino acid

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Wet Leu Ile Pro Leu Gin Cys Leu Phe Ser Ser Asp Arg Met Leu Thr
1 5
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Leu Ala Leu Val Glu Ile Lys Leu Glu Asp Leu Gln Ser Gln Leu His
20 25
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Met Leu Trp Lys Cys Ser Gln Asn Ile Ala Arg Cys Leu Leu Leu Leu
1 15
                                                                                                                                                                                                                                                                                                   Met Thr Phe Ser Ser Leu Lys Leu Phe Val Leu Thr Cys Ile Ile Lys
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (2) INFORMATION FOR SEQ ID NO: 337:
                                                                                                                                                 Gln Arg Ser Leu Ser Ser Asn Leu Val His Val Leu Leu Gln Pro Ala $15$
                                                                                                                                                                                                                         Gly Leu Glu Arg Phe Ile Ile Leu Arg Glu Val Cys Asn Gln Glu Ile 20 \ \ 30
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Phe Leu Thr Fro Trp Gln Lys Gly Glu Lys Cys Val Leu Gly Trp Val
20 25
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   (2) INFORMATION FOR SEQ ID NO: 338:
Leu Tyr Ser Ile Lys Tyr Met Pro Pro Gln Lys Lys
65 70 75
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                                                                            Phe Lya Asp Val Leu Val Thr Glu Ile Ile Cys Leu Cys Met Cys
50 . .
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Trp Lys Ser Ile Pro Gly Pro Ser Pro Arg Asn Gln His Arg
35 40 45
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 336:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           (i) SEQUENCE CHARACTERISTICS:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                    (i) SEQUENCE CHARACTERISTICS:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 337:
                                                                                                                                                                                                                                                                                                                                                                        X.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Phe Leu Ser Glu Ile Ser Xaa
35 40
                                                                                                                                                                                                                                                                                                                                                                    SEQUENCE DESCRIPTION: SEQ ID NO: 338:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (D) TOPOLOGY: linear
                                                                                                                                                                                                                                                                                                                                                                                            (A) LENGTH: 76 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     (D) TOPOLOGY: linear
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              (B) TYPE: amino acid
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    (A) LENGTH: 41 amino acids
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SEQUENCE DESCRIPTION: SEQ ID NO: 341:

(A) LENGTH: 26 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Arg Leu Phe Phe Ile Gly Phe Leu Leu Leu Phe Ser Phe Gly Leu $10 \ 15$

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(1) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (A) LENGTH: 26 amino acids (D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 342:

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SEQUENCE DESCRIPTION: SEQ ID NO: 342:

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Leu

Arg Gln Pro Ser Leu Ser Ala Glu His 20 25

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(2) INFORMATION FOR SEQ ID NO: 341:

(i) SEQUENCE CHARACTERISTICS:

30

Val Leu Val Gly Leu Val Ile Val Ile Val Ala Thr Glu Leu Met Val
20
25

Met Ala Lys Ile Ser Pro Phe Glu Val Val Lys Arg Thr Ser Val Pro 1 15

(A) LENGTH: 42 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 340:

Pro Gly Thr Ala Ala Ala Val Thr Gly Lys 35 40

25

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(1) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 340:

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Gly Leu Thr Arg Tyr Met Pro Pro Xaa Ser Xaa Leu Asn 20 25

Ser Glu

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Lys Val Tyr Ile Phe Leu Ile Phe Met Val Leu Ile Leu Pro Ser Leu 1 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 339:

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(2) INFORMATION FOR SEQ ID NO: 339:

(i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(D) TOPOLOGY: linear (A) LENGTH: 31 amino acids

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Thr Glu 25 Ŀÿ3 Asp Leu Ala Lys 20 Arg Ser Ser

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INFORMATION FOR SEQ ID NO: 343: 3 2 SEQUENCE CHARACTERISTICS:

(A) LENGTH: 157 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 343: ž.

2

Phe Ser Ser Leu Glu Phe Tyr Gln Lys Lys Lys Ser Arg Trp Pro

Asp Glu Cys Ile Pro Trp Glu Val Trp Thr Val Lys Val His Val Val 20 20

8

Ala Leu Ala Thr Glu Glu Glu Arg Gln Ile Cys Arg Glu Lys Val Gly
35
40

Glu Lys Leu Cys Glu Lys Ile Ile Asn Ile Val Glu Val Met Asn Arg 50

23

His Glu Tyr Leu Pro Lys Met Pro Thr Gln Ser Glu Val Asp Asn Val 65 8

Leu Arg Asp Val Gln Pro Tyr Leu Tyr Lys Ile Ser 85 90 95 Phe Asp Thr Gly

Thr Ser Val Thr Thr Thr Met Arg 105 Phe Gln lle Thr Asp Ala Leu Gly 100 33

Arg Leu lie Lys Asp Thr Leu Pro Ser Glu Arg Arg Trp lie Ser Gly 115 \$ Ser Leu Met Ala Pro Arg Pro Trp Leu Leu Gly Ile Ala Leu Leu 130 Ser

Glu Pro Ala Leu Gly His Trp Xaa 150 Leu Trp Ala Leu Gly 145 5

INFORMATION FOR SEQ ID NO: 344: 2 S

(A) LENGTH: 520 amino acids 3

(B) TYPE: amino acid
(D) TOPOLGGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 344:

55

Leu Pro Leu Pro Ala Ala Gly Arg Val Val Val S Phe Leu Met

Leu Ala Val Arg Arg Phe Gly Ser Arg Ser Leu Ser Thr Ala Asp Met 8

Leu Lys Ala 80 Glu Lys Glu Asp Asp Asp Lys Leu Leu Ala 60 Leu His Gln Asp Phe Pro Ser Val 90 95 8 Pro Pro Thr Lys Gly Leu Val Leu Gly Ile Tyr Ser Lys Ser Ala Gly Glu Asn Phe 55 Leu Asn Ile Ser Gly 23 Thr Phe Tyr Gly 85 컱유 Val Pro Gln Phe Thr Gly Lys Leu Arg Glu 65 Gly Lys Thr Arg 2

Leu Val Gly Leu Gly Lys Lys Ala Ala Gly Ile Asp Glu Glu Glu 100 100 Asn Trp His Glu Gly Lys Glu Asn Ile Arg Ala Ala Val Ala Ala Gly 125 Val 15

Pro Asp Ser Val Glu Val Ser 3 Arg Gln Ile Gln Asp Leu Glu 130 ξ

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Glu Tyr Asp Asp Leu Lys Gln Lys Lys Lys Met Ala Val Ser Ala Lys 170 Gly Asp Ala Gln Ala Ala Ala Glu Gly Ala Val Leu Gly Leu Tyr 150 \$ 5 25

Leu Tyr Gly Ser Gly Asp Gln Glu Ala Trp Gln Lys Gly Val Leu Phe 180 3

Gly Gln Asn Leu Ala Arg Gln Leu Met Glu Thr Pro Ala Asn 195 Met Thr Pro Thr Arg Phe Ala Glu Ile Ile Glu Lys Asn Leu Lys 210 Ala Ser g 35

11e 240 Thr Glu Val His Ile Arg Pro Lys Ser Trp 230 235 Ala Ser Ser Lys 8

Glu Glu Gln Ala Met Gly Ser Phe Leu Ser Val Ala Lys Gly Ser Asp 255

Glu

5

Pro Pro Val Phe Leu Glu Ile His Tyr Lys Gly Ser Pro Asn Ala 260 270 Ser Asn.Glu Pro Pro Leu Val Phe Val Gly Lys Gly Ile Thr Phe Asp 275 20

Met Gly Gly Ala Ala Thr Ile Cys Ser Ala Ile Val Ser Ala Ala 310 315 Leu Met Arg Ala Gly Ile Ser Ile Lys Ala Ser Ala Asn Met Asp 290 Asp 305 gy 55

Asn Leu Pro Ile Asn Ile Ile Gly Leu Ala Pro Leu Cys Glu 335 2 Ę,

Asn Met Pro Ser Gly Lys Ala Asn Lys Pro Gly Asp Val Val Arg Ala

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(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 234-amino acids

2 INFORMATION FOR SEQ ID NO: 346: Gln Ser Leu Arg Leu Asn Ala ,35

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Gly Lys Asp Ser Ile Asp Ile Asp Ile Ser Ser Arg Arg Arg Glu Asp 20 25

Thr Ile Leu Phe Leu Phe Leu Gln Leu Ser Ala Leu Arg Leu Ile Val 1 5

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SEQUENCE DESCRIPTION: SEQ ID NO: 345: ΘĐ (D) TOPOLOGY: linear TYPE: amino acid LENGTH: 39 amino acids

3 SEQUENCE CHARACTERISTICS:

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2 INFORMATION FOR SEQ ID NO: 345:

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Tyr Lau Ser Phe Gly Asp Phe Ala Phe Cys Ala Glu Leu Met Ile Gln 195 200 205

Val Phe Phe Ala Asn Asn Arg Phe Glu Thr Gly Lys Lys Lys Leu Gln 180 185

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Asn

Trp Thr Leu Gly Pro Val Asp Ser Gln Met Asp Map Met Asp Met 210 215

Asp Leu Asp Arg Asn Phe Ser Arg Thr Xaa 225 230

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Gln His Phe Leu Leu Ser Asp Arg Leu Ala Arg Asp Tyr Ala Ala Ile 165 170 175

35

βī Phe Ser Gln Asp Asn Ala Xaa 515

30 Lys Gly Met Thr Gly Arg Pro Thr Arg Thr Leu Ile Glu Phe Leu Leu 505

Asp Ile Ala Gly Val Met Thr Asn Lys Asp Glu Val Pro Tyr Leu Arg 485 485

25 Ala Ala Phe Leu Lys Glu Phe Val Thr His Pro Lys Trp Ala His Leu 475 470

25

Ser Lys Cly Thr Lys Lys Asp 115

Leu Asp Asp Ile Ser Thr Lys Thr Gly 120 125

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Ile Thr Leu Lys Ser Cys Arg Arg Gln Phe Asp Asn Pho Lys Arg Val 130 140

Phe Lys Val Val Glu Glu Wet Arg Gly Ser Leu Val Asp Asn Ile Gln 145 150 150

20

Ile Phe Gin Ile Pro Pro Ser Arg Gin Ala Leu Leu Ile Giu Arg Tyr 85 90 95

Met Leu Glu Arg Leu Leu His Ala Pro Pro Lys Leu Leu His Gln Leu 65 70 75 80

Tyr Ala Phe Asp Glu Ala Phe Val Arg Glu Val Leu Gly Lys Lys Leu 100

Asp Val Asn Asn Ile Gly Lys Tyr Arg Ser Ala Gly Ala Cys Thr Ala 450 455

20

Pro Leu Phe Glu His Tyr Thr Arg Gln Val Val Asp Cys Gln Leu Ala 435 440 445

Lys Leu Phe Glu Ala Ser Ile Glu Thr Gly Asp Arg Val Trp Arg Met 420 425

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Ala Ala Val Leu Gin Ser Asp Thr Met Asp His Tyr Arg Thr Phe His 50 60

Val Ala Leu Arg Val Arg Ser Gly Ile Leu Glu Gln Thr Gly Ala Thr \$35\$

Glu Asp Val Tyr Arg Leu Trp Leu Asp Gly Tyr Ser Val Thr Asp Ala 20 25

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Met Thr Ser Glu Leu Asp Ile Phe Val Gly Asn Thr Thr Leu Ile Asp 1 .15

(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 346:

Gly Ser Gly Ala Thr Gly Val Phe Thr Asn Ser Ser Trp Leu Trp Asn 410

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Xam Ile Leu Asn Ala Ala Thr Leu Thr Gly Ala Met Asp Val Ala Leu 385 390 190 Ile Leu Ala Asp Ala Leu Cys Tyr Ala His Thr Phe Asn Pro Lys 370 380

Lys Asn Gly Lys Thr Ile Gln Val Asp Asn Thr Asp Ala Glu Gly Arg 355 345

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Met Ala Ala Ala Val Ala Gly Met Leu Arg Gly Gly Leu Leu Pro Gln

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INFORMATION FOR SEQ ID NO: 347:

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55 Ξ SEQUENCE CHARACTERISTICS:

(A) LENGTH: 169 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 347:

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Ala Gly Arg Leu Pro Thr Leu Gln. Thr Val Arg Tyr Gly Ser Lys Ala 20 2

Val Thr Arg His Arg Arg Val Met His Phe Gln Arg Gln Lys Leu Met 35

Ala Val Thr Glu Tyr Ile Pro Pro Lys Pro Ala Ile His Pro Ser Cys 50 60 2 Leu Pro Ser Pro Pro Pro Gln Glu Glu Ile Gly Leu Ile Arg 65

Leu Leu Arg Arg Glu Ile Ala Ala Val Phe Gln Asp Asn Arg Met Ile 85 13

Cys Gln Asn Val Ala Leu Ser Ala Glu Asp Lys Leu Leu Ile 100 Ala Val Ala

2

Pro Ala Ala Glu Thr Gln Asp Pro Asp Glu Gly Leu Pro Gln 115 Pro 22

Cys Pro Phe Leu Trp Gly Thr Thr Cys Cys Trp Ser Val Lys Ser Pro 160 145 Gly Pro Glu Ser Pro Ser Trp Arg Ile Pro Ser Thr Lys Ile Cys 130

Arg Ser Arg Arg Trp Tyr Gly Ser Xaa 165 ജ

(2) INFORMATION FOR SEQ ID NO: 348: 35

(A) LENGTH: 43 amino acids (1) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 348:

Thr Ala His Leu Ile Leu Leu Glu Thr Leu Ser Val Cys Leu Trp Leu 30 Lys Arg Ser Phe Leu Leu Pro Leu Leu Val Gly Phe Leu Asp 10 15 Æ 45

Leu Ile Asp Ser Arg Cys Val Met Ser 35 Pro Ser S

INFORMATION FOR SEQ ID NO: 349: 3 55

(A) LENGTH: 78 amino acids (1) SEQUENCE CHARACTERISTICS: TYPE: amino acid

SEQUENCE DESCRIPTION: SEQ ID NO: 349: (D) TOPOLOGY: linear (X

S

Met Lys Glu Gly Pro Pro Cys Lys Arg His His Tyr Tyr Gln Asn Cys

Phe Gly Glu Thr Aen Gln Ile His 25 Gly Ala Lys Leu Leu Val Ser Leu 20 Ś

Leu Leu Glu Thr Gln Val Gly Thr Glu Lys Gly Gly Glu Arg Ile Trp 35

Glu Glu Lys Trp Arg Ile Ser Ser Thr Val Leu Phe Ile Ser Val Asn 50 60

2

Ser Tyr Val Glu Gly Ser Val Leu Glu Ile Lys Leu Phe Tyr 65 $^{75}\,$ 2

(2) INFORMATION FOR SEQ ID NO: 350:

2

(i) SEQUENCE CHARACTERISTICS:

Met Ser Glu Ile Leu Ser Leu Leu Phe Cys Leu Leu Gly Pro Ala Leu 10 15 15 SEQUENCE DESCRIPTION: SEQ ID NO: 350: (xT

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Asp Glu Arg Arg Glu Glu Lys Asp 20 ജ

(2) INFORMATION FOR SEQ ID NO: 351: 35

(A) LENGTH: 274 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (1) SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 351:

6

Met Ser Ser Ala Gly Thr Ala Thr Pro Leu Glu Met Asp His Lys Leu 1 5 10 15 Thr Ser Gin Pro Gly Arg Pro Ser Phe Tyr Cys Asn Ser Arg His Ser 20 5

ile Val Gly Ser Ser His Gln Leu Gly Phe Trp Phe Ser His Leu Glu 35

S

Ser Ser Gly Leu Lys Val Phe Gln Val Ser Leu Pro Cys Glu Cys Val 50 60

Leu Pro Thr Arg Ile Ala Ser Val Val Leu Ser Leu Met Ser Leu 70 75 80 Leu Val Val Gly Gln Ala Pro Ala Trp Glu Gly Ser Leu Leu Arg Gly 85 90 g g 55

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Ser Leu Leu Cys Cys Leu Pro Trp Gln Cys Ser Gly Gly Gly Phe Pro 165 170 175

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20 Ala Thr Leu Ala Leu Thr Cys His Cys Asp Lys Val His Val Ala Gly 195 200 205

25

Asp Xaa Glu 2 225

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dat sth Arg Gly Met Thr Arg Leu Gly Arg Val Ser Leu Thr Ser Ser 260 265

35 Ile Xaa

(2) INFORMATION FOR SEQ ID NO: 352:

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Ě SEQUENCE CHARACTERISTICS:

(A) LENGTH: 47 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 352:

45

Met Ile Phe Thr Ser Val Thr Lys Gly Ile Leu Leu Ile Ala Leu Trp 1 5

8 Val Pro Leu Phe His Phe Met Leu Ile Asp Ser Ile Leu Gly Pro Ser $20 \ \ 25$

55 Arg Leu Leu Thr Asp Gly Val Pro Phe Asn Pro Trp His Val Xaa 35 40

(2) INFORMATION FOR SEQ ID NO: 353:

(i) SEQUENCE CHARACTERISTICS:

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Gly Gln Val Ser Gln Val Leu Pro Ala Leu Ser Leu Gly Leu Val Phe 130 135 Leu Val Val Asp Val Gly Glu Arg Ile Leu His Gly Gln Arg Glu Val 115 120 Arg Pro Ala Gly Gly Ala His Leu Cys Ala Met Xaa Val Ile Glu Gly 100 105

5 Leu Cys Gln Gly Thr Val Glu Lys Val Ser Gly Ala Ala His Cys Ser 145

Thr Xaa Arg Cys Ser Arg Pro Tyr Phe Ser Scr His Lys Gly Val Ala 180

Leu Gly Lys Asp Txp Ala Ile Glu Gln Arg Arg Thr Cys Glu Ser 210 220

XAA XAA Pro Phe Thr Leu Ala Gly Leu Val Leu Val Leu 230 235

Cys Gln Val Val Leu Val Trp Ile Pro Gln Leu Gly Asp Lys 250 255

30

30 Phe Ala Leu His 50 25

Val Thr Gln Ala Lys Trp Asn Ser Trp Pro Ser Arg Arg Asn Ala Gly
35 40 45

Glu Ala Xaa Val Arg Ser Ser Lys Lys Trp Ile Pro Lys Ala Leu Xaa 20 25 30

Met Ser Ile Ser Gly Thr Asp Gly Leu Ile Leu Leu Leu Val Gly Leu 1 5 10

20

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SEQUENCE DESCRIPTION: SEQ ID NO: 354:

(A) LENGTH: 52 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 354:

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SEQUENCE CHARACTERISTICS

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 353:

(A) LENGTH: 3 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

Met Lys Thr

(2) INFORMATION FOR SEQ ID NO: 355:

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(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 132 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Glu His Cys Leu Tyr His Ser Val His Gly Ile Asn Pro Tyr Ile 1 15 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 355:

8

His Lys Asn Thr His Pro Ser Ile Asn Ile Tyr Met Val Trp Asp Glu 20 25 30

5

Gln Val Asn Ser Phe Glu Arg Glu Phe Val Pro Phe Phe Phe Leu Ile 35 40 45

Ile Leu Leu Asn Cys Cys Gln Leu Ser Asn Lys Gln Thr Glu Lys Leu $$50\$

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Phe Gly Lys Thr Leu His Thr Pro Phe Leu Ser Ser Ala Leu Lys Tyr 65 70 75 80

Arg Leu Asn Thr His Ile Leu Pro Val Phe Ser Tyr Ser Asp Ser Ile 90 95

S

Leu Thr Cys His Leu Ile Leu Ala Ser Tyr Phe Ser His Val Tyr Leu 100 105

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Pro Val Thr Cys Ile Cys Tyr Leu Asn Arg Lys Lys Asn Ile Gln Lys 115

Lys Lys Asn Xaa 130

(2) INFORMATION FOR SEQ ID NO: 356: 9

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 204 amino acids TYPE: amino acid

Gly Ser Arg Asp His Leu Phe Lys Val Leu Val Val Gly Asp Ala Æ

SEQUENCE DESCRIPTION: SEQ ID NO: 356:

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2

먊 Gln Asp Tyr Ser Gln Asp Ser 25 Leu Val Ser 뀵 2 2 2 2 2 Ala Val Gly

2

Thr Val Gly Val Asp Phe Ala Leu Lys Val 40 Lys His Tyr Lys Ser 35 Ser 23

Trp Asp Leu Gln Leu Glu Ile Val Arg ጟ Ş. Gln Trp Ser

Leu Tyr Tyr Arg Asp 80 Ala Gly Glu Arg Phe Thr Ser Met Thr Arg 65 70 75 2

Thr Phe 뵱 Phe Asp Val Thr Asn Ala 90 Ala Ser Ala Cys Val Ile Met

Lys Leu Thr Leu 110 Ser Asn Ser Gln Arg Trp Lys Gln Asp Leu Asp 105 Ser

35

Pro Asn Gly Glu Pro Val Pro Cys Leu Leu Leu Ala Asn Lys Cys Asp 115 Ser Pro Trp Ala Val Ser Arg Asp Gin Ile Asp Arg Phe Ser Lys 130 Ę 6

Glu Asn Gly Phe Thr Gly Trp Thr Glu Thr Ser Val Lys Glu Asn Lys 145 5

Asn Ile Asn Glu Ala Met Arg Val Leu Ile Glu Lys Met Met Arg Asn 175 20

Leu Ser Thr Gln Gly Asp Tyr Ile Asn 185 Glu Asp Ile Met Ser 180 퉑 Ser

Trp Ser Cys Cys Xaa 200 Leu Gln Thr Lys Ser Ser Ser 195 55

(2) INFORMATION FOR SEQ ID NO: 357:

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200

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SEQUENCE CHARACTERISTICS: 3

(A) LENCTH: 47 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 357:

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Met 11e Ser Leu I1e Phe Gln Leu Glu Glu Lys Leu Val Glu Lys $1 \ \ \, 1$

Leu Phe Phe Leu Lys Lys Gly Ser Gln Gly Ser Phe 8 Phe Phe Phe 9

Leu Arg Gly Val Pro Arg His Met Arg Val Lys Ile Val Asn Leu

13

(2) INFORMATION FOR SEQ ID NO: 358

(i) SEQUENCE CHARACTERISTICS:

2

(A) LENGTH: 73 amino acids

TYPE: amino acid TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 358: ž

23

Met Thr Tyr Val Thr Cys Leu His Val Cys Leu Leu Val Glu Phe Leu

Asn Ser Gln Leu Thr Asn His Arg Lys Tyr Tyr Phe Leu Ser Tyr Gly 20 25

2

Phe Trp Phe Thr Gly Leu Arg Gly Phe Ser Glu Tyr Leu Trp Pro Gln 15

Gln His Thr Ser Phe His Pro Asn Arg Asn Glu Ile Asn Phe Val Ser S0 60 35

Thr Asp Asn Arg Ile Trp Val Thr Xaa 65

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(2) INFORMATION FOR SEQ ID NO: 359:

(A) LENGTH: 102 amino acids (i) SEQUENCE CHARACTERISTICS

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(B) TYPE: amino acid

(D) TOPOLOGY: linear

Met Ser Asp Gln Glu Ala Lys Pro Ser Thr Glu Asp Leu Gly Asp Lys 1 5 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 359.

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Lys Glu Gly Glu Tyr Ile Lys Leu Lys Val Ile Gly Gln Asp Ser Ser 25 20

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Lys Glu Ile His Phe Lys Val Lys Met Thr Thr His Leu Lys Lys Leu 15 45 Glu Ser Tyr Cys Gln Arg Gln Gly Val Pro Met Asn Ser Leu Arg Phe

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Gly His Ser Thr Val Xaa 100

Gly Met Glu Glu Glu Asp Val Ile Glu Val Tyr Gln Glu Gln Thr Gly 95

Leu Phe Glu Gly Gln Arg Ile Ala Asp Asn His Thr Pro Lys Glu Leu 65 70 75 80

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Gln Val Leu Glu Ala Pro Gly Val Tyr Val Phe Gly Glu Leu Leu Asp \$35\$Met Ser Ala Glu Val Lys Val Thr Gly Gln Asn Gln Glu Gln Phe Leu 1 15 ... Leu Leu leu Val Gly Leu Val Tyr Leu Val Ser His Leu Ser Gln Arg 35 40 45 Thr Ser Met Pro Așn Val Arg Glu Leu Ala Glu Ser Asp Phe Ala Ser Thr Phe 50 55 ē Glu Ala Arg Asn Leu Pro Pro Leu Thr Glu Ala Gln Lys Asn Lys Leu 95 Arg 65 (2) INFORMATION FOR SEQ ID NO: 361: Leu Leu Thr Val Phe Ala Tyr Gly Thr Tyr Ala Asp Tyr Leu Ala 70 75 Leu Ala Lys Ser Ala Lys Gly Ala Ala Leu Ala Thr Leu Ile His ξ Ê 11e SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 361: Leu Leu Phe Leu Leu Met Met Leu Gly Val Arg Gly $20 \ 25 \ 30$ (A) LENGTH: 179 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

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20 6 330 25 15 Met Gly Phe (2) INFORMATION FOR SEQ ID NO: 360: Ē $\widetilde{\mathbf{x}}$ SEQUENCE CHARACTERISTICS: Pro Gln Trp His Leu Gly Asn His Ala Val Glu Pro Val 5 SEQUENCE DESCRIPTION: SEQ ID NO: 360: (B) TYPE: amino acid
(D) TOPOLOGY: linear LENGTH: 48 amino acids

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Asn His Xaa

Asp Ile Gin Arg Gin Asp Leu Ser Ala Ile Ala Arg Thr Leu Xaa Lys 165 170

Leu Asp Gln Arg Asn Gln Arg Leu Glu Val Asp Tyr Ser Ile Gly Arg 145 150 150

Tyr Ala Val Leu Leu Glu Ala Leu Ala Leu Arg Asn Val Arg Gln Leu 115 120 125

Arg His Leu Ser Val Val Thr Leu Ala Ala Lys Val Lys Cys Ile Pro 100 105 110

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2 INFORMATION FOR SEQ ID NO: 362:

(i) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 25 amino acids(B) TYPE: amino acid (D) TOPOLOGY: linear

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 362:

Met Lys Ser Ser Ser Leu Phe Phe Phe Phe Leu Ala His Phe Ile His 1 10 15

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(2) INFORMATION FOR SEQ ID NO: 363:

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Ser His Asp Leu Pro Gly Leu Cys Arg 20 25

Ξ SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 224 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 363:

3 Met Lys Phe Ala Ala Ser Gly Xaa Phe Leu His His Met Ala Gly Leu 1 15

8 Ser Ser Ser Lys Leu Ser Met Ser Lys Ala Leu Pro Leu Thr Lys Val 20 25 30

Val Gln Asn Asp 35 Ala Tyr Thr Ala Pro Ala Leu Pro Ser Ser Ile Arg 40 45

Thr Lys Ala Leu Thr Asn Met Ser Arg Thr Leu Val Asn Lys Glu Glu 50 50 Pro Pro Lys Glu Leu Pro Ala Ala Glu Pro Val Leu Ser Pro Leu Glu 65 70 75 80

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Met Lys Thr 175 Phe Pro Ser 270 Leu Gln Glu Leu Leu Ser Lys Cys Arg Thr Cys. 120 ĘŻ Ser 160 Glu Thr Arg Glu Val Leu Thr Pro Thr Ser Thr Ser Asp Asn Glu Thr 180 Ser Ser Ser Asm Ile Arg Ser Gln His Ala Glu Glu Gln 210 Ser Asn Asn Gly Arg Tyr Asp Asp Cys Lys Glu Phe Lys Asp Leu His 245 Thr Ser lie Ser Ala Val Leu Ser Asp Leu Ala Asp Leu Arg Ser Cys 275 Gly Gln Ala Leu Pro Ser Gln Asp Pro Glu Val Ala Leu Ser Leu 290 Ser Cys Gly His Ser Arg Gly Leu Phe Ser His Met Gln Gln His Asp 320 Leu Cys Arg Thr Ile Glu Ser Thr Ile His Val Val 325 Met Leu His Gln Asp His Ile Thr Phe Ala Met Leu Leu Ala Arg Ile 1 5 10 Lys Leu Lys Gly Thr Val Gly Glu Pro Thr Tyr Asp Ala Glu Phe Gln Leu Gin Giu Gin Giu Ala Lys Giu Arg 135 Ser Ser lle lle Asp Pro Gly Thr Glu Gln Asp Leu Pro 195 Pro Glu Asn Ser Ser Val Lys Glu Tyr Arg Met Glu Val Pro Ser 210 210 Lys Asp Asp Glu Gly Ala Thr Pro Ile Lys Arg Arg Val 150 Ser Asp Glu Glu His Thr Val Asp Ser Cys Ile Ser Asp 165 Cys Ser Lys Asp Ser Thr Leu Ala Glu Glu Glu Ser Glu 260 Thr Arg Ile Ser Gly Lys Gly Asn Gln Ala Ala Ser Xaa 340 SEQUENCE DESCRIPTION: SEQ ID NO: 365: (A) LENGTH: 467 amino acids (i) SEQUENCE CHARACTERISTICS: TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 365: Ser Ala Leu Ile Pro Thr Ę Ser Glu Asp Met Gln Gln Arg 1 130 Ile Leu Asp Thr Œ, Arg Asp Phe 225 Ę 1 E Asp 9 2 ೫ 35 5 8 23 2 6 S 55 Asn Pro Pro 190 Ser Asn 80 Leu Arg 95 Ser Asp 160 Leu Leu Leu Ser Val His Thr Pro Lys Gln Leu Asn Pro 100 Asp Ile Val Glu Leu Phe Cys Val Cys Gly Ala Leu Lys Arg Ala Arg 110 Phe Val Lys Lys Asp Asp 125 Ala Ile Thr Ala Tyr Lys Lys Tyr Asn Asn Arg Cys Leu Asp Gly Gln 130 Gin Pro Ile Leu Leu Arg Leu Ser Asp Ser Pro Ser Met Lys Lys Glu 175 Gln Pro Thr Glu Phe Lys Ile Lys Leu Xaa 215 Phe Ala Phe Ser Glu Ala Asn Cys Ala Asn Leu Ile Ser Thr Leu Ile 55 Gly Thr Lys Met Thr Val Asn Asn Leu His Pro Arg Val Thr Glu Glu Ala Glu Val Asp Pro Asp Thr Ile Leu Lys Ala Leu Phe Lys Ser Ser 205 Glu Tyr Ile Lys Cys Ile Leu Met Asp Glu Arg Thr Phe Leu Asn Asn 20 Thr His Phe Leu Leu Lys Val Gln Ser 40 Met Ser Lys Asn Cys lle Lys Leu Leu Cys Glu Asp Pro Val 1 5 Gln Tyr Gln Asn Leu Gln Ser Asp Phe 70 Arg Val Glu Ile Ser Lys Ala Ser Ala Ser Leu Asn Gly Asp 90 캶 Leu His Met Asn Gly Asn Val Ile 150 Asn Ser Ala Ser Ser Ser 185 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 364; Leu Val His Pro Gly Val Ala Glu Val Val 116 SEQUENCE CHARACTERISTICS: TYPE: amino acid INFORMATION FOR SEQ ID NO: 364: (D) TOPOLOGY: linear Zet Leu Pro Arg Arg Val TP. Phe Pro Met Lys Cys Asn Ala Ser Xaa Thr 210 Asn Ile Val Tyr Thr 35 Asn Leu Ile Ser 180 Ala Leu Ala 3 Gln Val Ser Glu 귶 દુ જ 3

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Glu Gln Phe Gly Ile Trp Leu Asp Ser Ser Ser Pro Glu Gln Thr Val 85 90 95 Ser Cys Leu Pro Ala 65 His Phe Leu Arg Gly Asn Glu Ile Val Leu Ser Ala Gly Ser Thr Pro 35 40 Ala Ser Gly His Val Glu Asp Leu Ala Ala Glu Gln Asn Thr Gln Ile 180 Ser Ile Met Glu Gln Pro Leu Asp Leu Thr His Ile Val Xaa Thr Glu 145 150 150 Ala Met Ala His Met Phe Val Ser Thr Asn Leu Gly Glu Ser Phe Met 130 $$13^\circ$$ Ile His Arg Leu Leu Leu Ile Gln Ala Phe Arg Pro Asp Arg Leu Leu 115 120 Val His Leu Ala Pro Gly Trp Leu Met Gln Leu Glu Lys Lys Leu His 225 230 230 235 Alm Ile Asn Thr Alm Val Lys Ser Gly Arg Trp Val Met Leu Lys Asn $210 \ \ \, 215$ Thr Ser lle Ala Ile Gly Ser Ala Glu Gly Phe Asn Gln Ala Asp Lys 195 200 205 Val Lys Pro Asn Thr Pro Val Leu Met Cys Ser Val Pro Gly Tyr Asp 165 170 Pro Tyr Leu Leu Tyr bhe Leu Leu Ala Trp Phe His Ala Ile Ile Gln Glu Arg Leu 305 310 Ser Ile Pro Val Ser Arg Ile Cys Lys Ser Pro Asn Glu Arg Ala Arg 290 295 Phe Glu Pro Pro Pro Gly Xaa Lys Ala Asn Met Leu 275 280 Asn Pro Lys Val Pro Val Asn Leu Leu Arg Ala Gly Arg Ile Phe Val 260 265Ser Leu Gln Pro His Ala Cys Phe Arg Leu Phe Leu Thr Met Glu Ile 245 250 250 Arg Tyr Ala Pro Leu Gly Trp Ser Lys Lys Tyr Glu Phe Gly Glu Ser 325 330 335 Trp Ser Glu Glu Thr Pro Ala Thr Pro Ile Gly Gln Ala 100 105 110 Phe Lys Asp Leu Ile Ala Lys Val Gln Ala Asp 70 75 80 Arg Thr Phe Ser 285

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INFORMATION FOR SEQ ID NO: 366:

(1) SEQUENCE CHARACTERISTICS

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Met Ala Asp Glu Ala Thr Arg Arg Val Val Ser Glu Ile Pro Val Leu 1 5 10

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 366:

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Asp Leu Arg Ser Xaa Cys Asp Thr Val Asp Thr Trp Leu Asp Asp Thr

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10 25 20 15 нів Asp Gln 465 Glu Phe Asp Gln Arg Leu Leu Asn Thr Phe Leu Glu Arg Leu Phe Thr 385 390 395 Leu Lys Thr Leu Met Ala Gln Ser Ile Tyr Gly Gly Arg Val Asp Asn 370 380 Ala Lys Gly Arg Gln Asn Ile Ser Pro Asp Lys Ile Pro Trp Ser Ala 355 360 365 Pro Ala Gln Gln Arg Arg Glu Ser Pro Pro Tyr His Thr Gly Cys Gly $450 \ \ \, 460$ Ala Val Gly Gly Val Ala Pro Arg His Pro Asp Ala Leu Leu Ala Gly 435 440 445 His Lys Asp Ile Gln Met Pro Asp Gly Met Gln Ala Arg Gly Val Cys $425 \ \ \,$ Thr Arg Ser Phe Asp Ser Glu Phe Lys Leu Ala Cys Lys Val Asp Gly
410
415

Lys Thr Asn Ala Gly Pro Arg Asp Arg Glu Leu Trp Val Gln Arg Leu $20 \ 30$ Ala Asp Asn Asp Trp Phe Arg Leu Glu Ser Asn Lys Glu Gly Thr Arg 50 55 Lys Glu Glu Tyr Gln Ser Leu Ile Arg Tyr Val Glu Asn Asn Lys Asn 45

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Trp Phe Gly Lys Cys Trp Tyr Ile His Asp Leu Leu Lys Tyr Glu Phe 65 70 75 80 Asp Ile Glu Phe Asp Ile Pro Ile Thr . 85 Ile Ala Val Pro Glu Leu Asp Gly Lys Thr Ala Lys Met Tyr Arg Gly
105 106 Tyr Pro Thr Thr Ala Pro Glu 90 95

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Gly Lys Ile Cys Leu Thr Asp His Phe Lys Pro Leu Trp Gly Gln Glu 115 120 125

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Asp 320.

Lys

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Leu ile Glu Arg Gln Lys Lys Val Lys Leu Phe Cys Leu Glu Thr Phe. 255 Ala Arg Pro Trp Ala Arg Val Ile Gly Leu Arg Ile Trp Pro Gln 70 70 Leu Ser Asp Leu lle Ala His Gin Lys Gly Gln Ile Glu Lys Gln Pro 260 270 Leu Glu Arg Lys Leu Ile Leu Val Gln Val Ile Pro Val Val Ala 290 Ser Val Arg Leu Gln Ile Ser Thr Pro Asp Ile Lys Asp Asn 335 ile Val Ala Gin Leu Lys Gin Leu Tyr Arg ile Leu Gin Thr Gin Giu 345 Ser Trp Gln Pro Met Gln Pro Thr Pro Ser Met Gln Leu Pro Pro Ala 360 Met Gly Ser Ser Val Leu Pro Phe Cys Val Cys Val Thr Ser Pro Ser Leu Gly Gly Arg Cys Ile Gln Gly Arg Phe Ala Ser His Ser Lys Phe 20 20 10 Phe Gly Arg Lys Thr Ala Ser Phe Gly Ala Val Gly Glu Thr 35 Pro Pro Asp Gln Glu Pro Gln Lys Glu Thr Glu Pro Ala Thr Ser Ser 50 60 Glu Ile Tyr Leu Cys Phe Gly Glu Glu Trp Pro Asp Gly 275 $$280\$ $\mbox{Arg Het}$ lle Tyr Glu Met Phe Ser Gly \mbox{Asp} Phe Thr $\mbox{Arg Ser}$ Phe 305 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 368: SEQUENCE CHARACTERISTICS:
(A) LENGTH: 83 amino acids INFORMATION FOR SEQ ID NO: 369: (2) INFORMATION FOR SEQ ID NO: 368: (D) TOPOLOGY: linear Pro Pro Gln Xaa 370 Pro Asn Xaa Trp Gly Pro Phe Ser Gly His 65 Pro 3 3 S 2 2 ន 22 2 35 6 45 S 55 8 His Gly Ser Gly Ala Gly Ser 140 Tyr Asp Gly Thr Lys Glu Val Pro Met Asn Pro Val Lys Ile Tyr 5 10 Phe Leu Asn Ile Asn Gly Ser Pro Met Ala Pro 70 80 Asp Leu Asp Ile Lys Phe Gln Tyr Arg Gly Lys Glu Tyr Gly Gln 130 Pro Val Xaa Leu 175 Lys Leu Leu Asp Val Met Asp Arg Gly Leu Ile 200 Lys Val Tyr Trp Ser Gly Pro Cys Ala Pro Ser Leu Val Ala Pro Asn 215 Gin Val Cys Asp Ile Pro Gin Pro Gin Gly Ser Ile Ile Asn Pro Gly 20 20 Ser Thr Gly Ser Ala Pro Trp Asp Glu Lys Asp Asn Asp Val Asp Glu 35 Glu Asp Glu Glu Asp Glu Leu Asp Gln Ser Gln His His Val Pro Ile 50 60 Ala Ser Val Gly Asn Cys Ser Val Gly Asn Cys Ser Pro Glu Ala Val 85 Trp Pro Lys Thr Glu Pro Leu Glu Met Glu Val Pro Gln Ala Pro Ile 100 Phe Tyr Ser Ser Pro Glu Leu Trp Ile Ser Ser Leu Pro Met 115 Asn Pro Gln Gly Cys Arg Leu Phe Tyr Gly Asp 150 Asn Glu Lys Gln 190 Leu Glu Val Ser Gly His Ala Ile Tyr Ala Ile Arg Leu Cys Gln Cys 210 SEQUENCE DESCRIPTION: SEQ ID NO: 367 Leu Gly Pro Met Pro Asp Gln Glu Glu Leu Phe Gly 165 Glu Gln Val Lys Phe Pro Gly Pro Glu His Ile Thr 180 (A) LENGTH: 373 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear Ser INFORMATION FOR SEQ ID NO: 367 7 Asn Pro Xaa 150 Cys Ala Gln Ile Trp Thr Ser Ser 130 Phe Thr Ser i 195 Gln Asp Thr Phe Pro 65 Met Xea Gly Ser Gly 145 Ser Met Thr Val (x;) Gln Pro Lys Leu 3 த 14 5 2 15 2 25 30 35 6 45 S 55 S

575

S 3 55 Gly Thr Glu Ile Gly Val Leu Ala Lys Ala Phe Ile Asp Gln Gly Lys $50 \ \ 55$ Leu Lys His Leu Ser Ser Gly Asp Leu Leu Arg Asp Asn Met Leu Arg 35Gly Ser Gly Lys Gly Thr Val Ser Ser Arg Ile Thr Thr His Phe Glu $20 \ \ 30$ Met Gly Ala Ser Ala Arg Leu Leu Arg Ala Val Ile Met Gly Ala Pro 1 15 2 Tyr Trp Thr Met Xaa 20 Met Leu Leu Ser Val Ala Ile Phe Ile Leu Leu Thr Leu Val Tyr Ala 10 Leu Ile Pro Asp Asp Val Met Thr Arg Leu Ala Leu His Glu Leu Lys $65 \hspace{1.5cm} 70 \hspace{1.5cm} 75$ Pro Lys Thr Val Gly Ile Asp Asp Leu Thr Gly Glu Pro Leu Ile Gln 145 150 Pro Gin Ala Glu Ala Leu Asp Arg Ala Tyr Gin Ile Asp Thr Val Ile 100 105 Trp Ile His Pro Ala Ser Gly Arg Val Tyr Asn Ile Glu Phe Asn Pro . 130 Asn Leu Asn Val Pro Phe Glu Val Ile Lys Gln Arg Leu Thr Ala Arg 115 120 125 Asn Leu Thr Gln Tyr Ser Trp Leu Leu Asp Gly Phe Pro Arg Thr Leu 95 Glu Asp Gln Arg Glu Asp Asp Lys Pro Glu Thr Val Ile Lys Arg Leu Lys Ala Tyr 165 170 175 Leu Glu Thr Phe Ser Gly Thr Glu Thr Asn Lys Ile Trp Pro Tyr Val 195 200 205 INFORMATION FOR SEQ ID NO: 370: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 370: (XX) E (i) SEQUENCE CHARACTERISTICS SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: 369: The Lys Pro Val Leu Glu Tyx Tyx Gln Lys Lys Gly Val 180 185 (A) LENGTH: 21 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 227 amino acids (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 372:

(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENOTH: 51 amino acids

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Met Arg Ala Val Phe Pro Cys Cys Pro Phe Leu Thr Leu Met Leu Pro
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Leu Glu Cys Leu Val Gly Met Ile Met Cys Tyr Leu Gly Ile Ser 20

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Phe Thr Asp Thr Arg Lys Thr Ala Gly Leu Lys Lys Lys Lys Lys Lys Lys 45

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(2) INFORMATION FOR SEQ ID NO: 372:

(i) SEQUENCE CHARACTERISTICS:

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25 20 2 5 30 S Val Thr Pro 225 Tyr Ala Phe Leu Gln Thr Lys Val Pro Gln Arg Ser Gln Lys Ala Ser 210 215 220 Met Phe Leu Asn Cys Glu Ile Leu Glu Tyr Cys Tyr Tyr Leu Thr Gln 1 15 Lys Thr Ala Lys Gly Gly Gly Gln Glu Ala Leu Thr Cys Thr Xaa 65 70 75 Trp Ala Ile Lys Ala Gin Leu Lys Ile Giu Asn Lys Asp Leu Asp Asn 50 60 Lys Ile Ile Arg Cys Ser Ile Thr Ala Val Ser Asp Ser Ser Thr Ser 35 40 45 . Leu Lys Ile Ser Met Gly Lys Tyr Leu Ser Ile Pro Thr Val Leu Leu 20 25 (2) INFORMATION FOR SEQ ID NO: 371: Ξ X. SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 371: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 79 amino acids

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2 INFORMATION FOR SEQ ID NO: 373:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 61 amino acids

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SEQUENCE DESCRIPTION: SEQ ID NO: 373: (B) TYPE: amino acid (D) TOPOLOGY: linear Ī Phe Leu Met Arg Met His Leu Cys Phe Cys Lys Tyr Cys Cys Ser 10 ٨

Tyr Leu Trp 30 Phe Ile Val Thr Pro Thr Ser Thr Ser Asn Thr Ala Ser 20

2

Ser Ser Trp Ala Cys Pro Trp Ile Ser Ala Ser Met Ala Gly Arg Gly

Glu Xaa 9 Asn Ala Val Thr Arg Glu Gly Leu Pro 55 3 8 뀹

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(2) INFORMATION FOR SEQ ID NO: 374:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 40 amino acids

TYPE: amino acid (D) TOPOLOGY: Linear

SEQUENCE DESCRIPTION: SEQ ID NO: 374: ž.

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Leu Aan Thr His Thr Leu Cys Phe Val Leu Phe Cys Phe 5 Met Ser Leu

Ile Asn Glu Glu Lys Leu Ala Asn His Leu Ala Phe Arg 20 30 Ser Ē 뵱 8

Phe Ile Val Phe Xaa ile Leu Phe 6 35

35

INFORMATION FOR SEQ ID NO: 375:

2

(A) LENGTH: 44 amino acids SEQUENCE CHARACTERISTICS: 6

TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 375:

Met Cys Ser Gly Gln Ser Gln Val Trp Lys Met Ala Leu Gln Ala Leu 45

Asp Ser Glu Thr Val Val Ile Leu Pro Asp Met His Leu Ile Leu Ser 20 S

Leu Ile His Asn Ala Arg Pro Cys Leu Xaa 35 Leu Arg

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(2) INFORMATION FOR SEQ ID NO: 376:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 203 amino acids

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(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 376:

Met Leu Ile Ser Glu Glu Glu Ile Pro Phe Lys Asp Asp Pro Arg Asp. 1 5

Glu Thr Tyr Lys Pro His Leu Glu Arg Glu Thr Pro Lys Pro Arg Arg 20

Lys Ser Gly Lys Val Lys Glu Glu Lys Glu Lys Lys Glu Ile Lys Val 35 46

2

Ser 80 Glu Val Glu Val Glu Val Lys Glu Glu Glu Asn Glu Ile Arg Glu Asp 50 60 Ş Glu Glu Pro Pro Arg Lys Arg Gly Arg Arg Arg Lys Asp Asp 65 75

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Pro Arg Leu Fro Lys Arg Arg Lys Lys Pro Pro Ile Gln Tyr Val Arg 85

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Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala His Pro Arg Tyr Leu 100

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Gln His His Ile Lys Tyr Gln His Leu Leu Lys Lys Lys Tyr Val Cys 115

Pro His Pro Ser Cys Gly Arg Leu Phe Arg Leu Gln Lys Gln Leu Leu 130 Arg His Ala Lys His His Thr Asp Gln Arg Asp Tyr Ile Cys Glu Tyr 145 160 8

Ser His Asn Leu Ala Val His Arg Met 170 175 Cys Ala Arg Ala Phe Lys Ser 35

Ile His Thr Gly Glu Lys His Tyr Asn Val Arg Ser Val Asp Leu Leu 180

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Val Asp Lys Arg His Leu Leu Ile Gly Thr Xea 195

(2) INFORMATION FOR SEQ ID NO: 377:

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(A) LENGTH: 29 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 377: (x

Met Leu Pro Arg Arg Thr Phe Tyr Phe 1yr Phe 11e Phe 11e Phe 11 15 $$\rm 1$

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Leu Ala Ser Phe Trp Gly Phe Thr Leu Arg Ala Ser Phe 20

Met Phe Asp Ser Leu Ser Tyr Phe Lys Gly Ser Ser Leu Leu Leu Met 1 5 10 Phe Tyr Asn His Lys Phe Leu Xaa 130 · 135 Phe Met Lys Lys Gly Lys Glu Leu Phe Ile Gln Gln Glu Arg Phe Phe Leu Asn 90 95 Ąsp Tyr Leu His Asn His Ser Tyr Ala Ser Ile Gln Ser Asp Asp Leu Trp $_{35}$ Leu Lys Thr Gly Leu Met Ala Trp Arg Arg Glu Pro Ala Ser Gly Leu Ala Ala Cys Trp
1 10 15 (2) INFORMATION FOR SEQ ID NO: 379: Met Lys Thr Trp Thr Leu Gln Lys Gly Phe Pro Leu Val Thr Val Gln 65 70 70 80 2 Ser Lys Pro Glu Ile Gln Pro Ser Asp Thr Arg Tyr Met Pro Ser Phe $100\,$ Ser Phe Asn Glu Val Thr Asn Gln Thr Leu Asp Val Lys Arg Met 50 \$50r Cys His Leu F 115 (X <u>X</u> Ξ Ser Ser Gly Ser Arg Pro Trp 35 40 SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: Tyr Leu Ser Glu Asp Val Phe Gln His Ala Val Val Leu 20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 378: SEQUENCE DESCRIPTION: SEQ ID NO: 379: (A) LENGTH: 41 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 136 amino acids Phe Cys Thr Leu Arg Trp Lys Tyr Phe Glu Val 120 125

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INFORMATION FOR SEQ ID NO: 378:

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INFORMATION FOR SEQ ID NO: 380:

(1) SEQUENCE CHARACTERISTICS:

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Glu Ala Lys Ala Asn Val Cys Cys Val Lys Phe Ser Pro Ser Ser Arg

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8 5 25 5 30 ઝ 45 6 55 8 Pro Gin Phe Glu Ala Pro Ser Pro Ser His Ser Ser Ile Ile Asp Ser 50 55 Glu Glu Met Ser Gly Leu Tyr Ser Pro Val Ser Glu Asp Ser Thr Val 35 Gln Ile Gln Lys Glu Leu Ser Val Leu Glu Glu Asp Ile Lys Arg Val 20 25 Met Glu Phe Leu Lys Val Ala Arg Arg Asn Lys Arg Glu Gln Leu Glu 1 15 Lys Gln Pro Trp Tyr Asn Ser Thr Leu Ala Ser Arg Arg Lys Arg Leu 90 95 The Glu Tyr Ser Gln Pro Pro Gly Phe Ser Gly Ser Ser Gln The $65\,$ Gln Glu Cys Leu Ser Lys 130 Ser Arg Ile Ser Asp Asp Ser Arg Thr Ala Ser Gin Leu Asp Glu Phe 115 120 125 Thr Ala His Phe Glu Asp Leu Glu 100 Ala Thr Leu Ser Tyr Ala Ser Asp Leu Tyr Asn Gly Ser Ser Ile Val 145 150 Ala Val Val Thr Lys Lys Ile Lys Val Tyr Glu Tyr Asp Thr Val Ile Gln Asp 180 Ser Ser Ile Glu Phe Asp Arg Asp Cys Asp Tyr Phe Ala Ile Ala Gly 175 Ser Asp Tyr Glu Gly Thr Val Ile Leu Trp Asp Gly Phe Thr Gly Gln 235 230 Ile Ser Cys Ile Ser Trp Ser Ser Tyr His Lys Asn Leu Leu Ala Ser 210 215 Lys Val Lys I 275 Arg Ser Lys Val Tyr Gln Glu His Glu Lys Arg Cys Trp Ser Val Asp 255 255 Phe Asn Leu Met Asp Pro Lys Leu Leu Ala Ser Gly Ser Asp Asp Ala 260 265 270 (xi) SEQUENCE DESCRIPTION: Asp Ile His Tyr Pro Glu Asn Glu Met Thr Cys Asn Ser Lys 195 205 Leu Trp Ser Thr Phe Thr Arg Tyr Asn Ser Val Arg Pro Leu 135 Asn Leu Asp Asn Ser Val Ala Ser Ile 280 285 Gln Cys Tyr Phe Ser Thr Arg 105 SEQ ID NO: 380: Mer Eys 80

(A) LENGTH: 468 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Tyr His Leu Ala Phe Gly Cys Ala Asp His Cys Val His Tyr Tyr Asp 310 315 Lys Gln Pro Ile Met Val Phe Lys Gly His Arg Lys 325 Ala Val Ser Tyr Ala Lys Phe Val Ser Gly Glu Glu Ile Val Ser Ala 340 8 295 Leu Arg Asn Thr 290 2

Leu Arg Ser Phe Lys Gly His Ile Asn Glu Lys Asn Phe Val Gly Leu 370 Ala Ser Asn Gly Asp Tyr Ile Ala Cys Gly Ser Glu Asn Asn Ser Leu 185 2 8

Lys Pro Tyr Cys 365

Asp Ser Gln Leu Lys Leu Trp Asn Val Gly 355

Ser Thr

Tyr Leu Tyr Tyr Lys Gly Leu Ser Lys Thr Leu Leu Thr Phe Lys Phe 410 Lys Ser Val Leu Asp Lys Asp Arg Lys Glu Asp Asp Thr 420 Asp Thr Val 23

Asn Glu Phe Val Ser Ala Val Cys Trp Arg Ala Leu Pro Asp Gly Glu 415

Asn Val Leu Ile Ala Ala Asn Ser Gln Gly Thr Ile Lys Val Leu 450 Leu Val Xaa Glu 465 35

Ser

8

(2) INFORMATION FOR SEQ ID NO: 381:

(A) LENOTH: 29 amino acids SEQUENCE CHARACTERISTICS: TYPE: amino acid Ξ

Met Arg Lys Glu Asp Gly Phe Trp Phe Phe Phe Phe Leu Phe Phe Phe l 10 $$\rm 15$ SEQUENCE DESCRIPTION: SEQ ID NO: 381: ž

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Val Val Gly Ser Lys Phe Val Asn Gly Asn Lys Leu Val 20 S

(2) INFORMATION FOR SEQ ID NO: 382:

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(A) LENCTH: 29 amino acida
(B) TYPE: amino acid
(D) TOPOLOCY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 382: (1) SEQUENCE CHARACTERISTICS:

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Met Pro Leu Ala Pro Tyr Cys Asp Leu Leu Val Ala Leu Ser Phe Ala 1 5 15

Ser Ser Asp Phe Thr Leu Val Leu Glu Ser Pro Val Asp

(2) INFORMATION FOR SEQ ID NO: 383: 2

SEQUENCE DESCRIPTION: SEQ ID NO: 383: (A) LENGTH: 138 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

15

Met Asn Ser Leu Val Ser Trp Gln Leu Leu Leu Phe Leu Cys Ala Thr. 1 5 15 15 (X

2

Pro Thr Gly Gln Gln Leu Glu Ser Leu Gly Leu Leu Ala Pro Gly Glu 35 His Phe Gly Glu Pro Leu Glu Lys Val Ala Ser Val Gly Asn Ser Arg \$20\$

Thr Glu Arg Lys Pro Ala Ala Thr Ala Arg Leu 55 Gln Ser Leu Pro Cys 50

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Ser Arg Arg Gly Thr Ser Leu Ser Pro Pro Pro Glu Ser Ser Gly Ser 65 75 80 Pro Gin Gin Pro Gly Leu Ser Ala Pro His Ser Arg Gin Ile Pro Ala 90 85 3

Pro Gln Gly Ala Val Leu Val Gln Arg Glu Lys Asp Leu Pro Asn Tyr 110

35

Asn Trp Asn Ser Phe Gly Leu Axg Phe Gly Lys Axg Glu Ala Ala Pro 125 8

Gly Asn His Gly Arg Ser Ala Gly Arg Gly 130

INFORMATION FOR SEQ ID NO: 384: 3

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(A) LENGTH: 74 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 384: (i) SEQUENCE CHARACTERISTICS:

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Met Ser Cys Phe Ile Asp Ser Xaa Asp Ser Lys Ile Leu His Leu Leu 1 5 10 15 55

Val Val Ser Phe lle Cys Xaa Leu Phe Leu Leu Ile Leu Thr His Gly 20 10

83

5 45 6 35 30 25 20 7 S 8 55 Arg Glu Asp Phe Phe Val Leu Pro Xaa Ala 65 (70 Agn Ile Leu Ile Leu Arg Xaa Phe Phe Ser Val Xaa Xaa His Ser Leu Lys 35 40 Met Ser Ala Gly Glu Val Glu Arg Leu Val Ser Glu Leu Ser Gly Gly 1 15 (2) INFORMATION FOR SEQ ID NO: 385: Val His Val Thr Ile Gly Asp Ile Lys 85 Glu Thr Glu Asp Asp Ser Asp Ser Asp 65 rle Glu Asp Glu Thr Ala Glu Asn Gly Val Pro Val Glu Arg Pro Glu Glu Glu Asn Ala Ser Ala Asn Pro Pro Ser Gly
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45 Phe Asn Tyr Gly Phe Asn Glu Asp Thr 165 Pro Gly Sex Ile Asn Gly Val Pro Leu Leu Glu Val Asp Leu Asp Sex 130 135 Val Tyr Gly Thr Thr Gly Thr Lys Val Lys Gly Val Asp Leu Asp Ala 115 Ser The Gly Gly Phe Glu Asp Lys Pro Trp Arg Lys Pro Gly Ala Asp Leu Ser Asp Tyr 145 150 Thr Asn Lys Ile Thr Val Gln Gln Gly Arg Thr Gly Asn Ser Glu Lys 195 200 Gln Lys Asn Leu Glu Glu Tyr Leu Ile Leu Met Asn Lys Ala Leu Leu Thr 50 55 Tyz Gly Thr Ala Pro Val Asn Leu Asn Ile Lys Thr Gly Gly Arg $100\,$ Ξ ξ Arg Ile Arg Met Gly Leu Glu Val Ile Pro Val Thr Ser Thr 180 185) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 521 amino a.

(B) TYPE: amino acid

(D) TOPOLOGY: linear Asp Glu Glu Glu Olu Trp Leu Tyr Gly Asp Glu Asn Glu 20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 385: Thr Gly Ala Pro Gln Tyx Gly 90 95 Ser Trp Lys Ala Tyr Cys Glu Lys 170 175 Asp Asp Asp Glu Asp Asp 75 Lys Pro Lys Val Thr 60

5 25 20 2 30 6 35 55 8 3 Gly Ala Pro Pro Thr His Leu Pro Pro Pro Pro Phe Leu Pro Pro Pro 290 295 300 Thr Glu Val Asp Asn Asn Phe Ser Lys Pro Pro Pro Phe Phe Pro Pro 275 280 285 $\mbox{Arg Ala Asn Glu Asn Ser Asn Ile Gln Val Leu Ser Glu Arg Ser Ala <math display="inline">260$ Asp Val Ile Gly Gin Thr Ile Thr Ile Ser Arg Val Glu Gly Arg Arg 245 250 255 Leu Phe Lys Thr Gly Leu Pro Pro Ser Arg Arg Leu Pro Gly Ala Ile 225 230 230 235 ř Pro Thr Val Ser Thr Ala Pro Pro Leu Ile Pro Pro Pro Gly Phe Pro 305 310 315 Gly Asn Val Ala Phe Pro His Leu Pro Gly Ser Ala Pro Ser Trp Pro 365 Gly His Ser Ser Gly Tyr Asp Ser Arg Ser Ala Arg Ala Phe Pro Tyr 340 345 Glu Lys Glu Glu Arg His Arg Glu Glu Glu Glu 450 455 Tyr Arg Glu Tyr Ala Glu Arg Gly Tyr Glu Arg His Arg Ala Ser Arg 435 Arg Asp Arg Glu Arg Glu Arg Thr Arg Glu Arg Glu Arg Glu Arg Asp 415 Lys Asp Arg Asp Arg Glu Arg Asp Arg Asp Arg Glu Arg Asp Arg Asp 385 390 Ser Glu Glu Gly Asp Ser His Arg Arg His Lys His Lys Lys Ser Lys Arg 485 490 495 Thr Arg His Lys Ser Ser Arg Ser Asn Ser Arg Arg Arg His Glu Ser 475 470 Ser Thr Glu Ala Thr Pro Ala Glu Xaa 515 520 Ser Ser Pro Thr Pro Ser Val Phe 420 Leu Val Asp Thr Ser Lys Gln Trp Asp Tyr Tyr Ala Arg Arg Glu 370 380 Pro Pro Gly Ala Pro Pro Pro Ser Leu Ile Pro Thr Ile Glu Ser 330 335 Lys Glu Gly Lys Glu Ala Gly Ser Glu 505 Asn Ser 425 Asp Glu Glu Arg Tyr Arg 430 Pro Ala Pro Glu Gln Glu 510

584

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Glu Thr Ala Leu|Pro Ser Thr Lys Ala Glu Phe Thr Ser Pro Pro Ser 210 215

INFORMATION FOR SEQ ID NO: 386: 3

SEQUENCE CHARACTERISTICS

(A) LENGTH: 137 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 386: ž. Met Asn Ser Arg Gly Ile Trp Leu Ala Tyr Ile Ile Leu Val Gly Leu 9

Leu His Met Val Leu Leu Ser Ile Pro Phe Phe Ser Ile Pro Val Val 25

Trp Thr Leu Thr Asn Val Ile His Asn Leu Ala Thr Tyr Val Phe Leu $40 \ \ 45$

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His Thr Val Lys Gly Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ala 50

Arg Leu Leu Thr His Trp Glu Gin Met Asp Tyr Gly Leu Gin Phe Thr 65 2

ş g S Ser Ser Arg Lys Phe Leu Ser Ile Ser Pro Ile Val Leu Tyr 90

52

Ala Ser Leu Leu Ser Val Leu Leu Pro Lys Leu Pro Gln Phe His Gly .115 ဓ္ဌ

Ala Ser Phe Tyr Thr Lys Tyr Asp Ala Ala His Phe Leu Ile Asn Thr 100

Val Arg Val Phe Gly Ile Asn Lys Tyr 130

35

(2) INFORMATION FOR SEQ ID NO: 387:

(A) LENGTH: 186 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS: £

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SEQUENCE DESCRIPTION: SEQ ID NO: 387:

<u>z</u>

Met Ala Ala Gln Lys Asp Gln Gln Lys Asp Ala Glu Ala Glu Gly Leu 1 5 10 15 45

Thr Thr Leu Leu Pro Lys Leu Ile Pro Ser Gly Ala Gly Arg 20 Glu Trp Leu Glu Arg Arg Arg Ala Thr Ile Arg Pro Trp Ser Thr Phe 45 46 Ser Gly S

phe Ser Arg Pro Arg Asn Leu Gly Glu Leu Cys 55 60 Asp Gln Gln Arg 50 Val 55

Gln Arg Leu Val Arg Asn Val Glu Tyr Tyr Gln Ser Asn Tyr Val Phe 65

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Val Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met Leu 95 $\,$ 95 $\,$ Leu Ala Val Phe Ghy Ala Cys Tyr Ile Leu Tyr Leu 100 110 Leu Val Ala

Arg Glu Val Ser Pro 125 Leu Glu Ser Lys Leu Val Leu Phe Gly 115 Arg Thr

Phe Pro Phe Phe Trp Ala His Gln Tyr Ala Leu Ala Gly Gly Ile Ser 130 2

Leu Ala Gly Ala Gly Ser Ala Val Phe Trp Val Leu Gly Ala Thr Leu 145

Val Val Ile Gly Ser His Ala Ala Phe His Gln Ile Glu Ala Val Asp 175

2

Gly Glu Glu Leu Gln Met Glu Pro Val Xaa 180 185 20

(2) INFORMATION FOR SEQ ID NO: 388:

23

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1 amino acids TYPE: amino acid (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 388:

33

Met 1

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(2) INFORMATION FOR SEQ ID NO: 389:

(i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 389:

Met Leu Ser 11e Phe Tyr Phe Ala 11e Pro Val Gly Ser Gly Leu Gly 15 10 10 15 15 45

Leu Arg Val Thr Pro Gly Leu Gly Val Val Ala Val Leu Leu Leu 35 Tyr ile Ala Gly Ser Lys Val Lys Asp Met Ala Gly Asp Trp His Trp 20 Ala

S

Leu Val Val Arg Glu Pro Pro Arg Gly Ala Val Glu Arg His Ser 50 60 먑

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Trp Ala Asp Leu Arg Ala Leu Asn Pro Thr Ser Trp 70 Pro Pro 85 65 Leu Ala Arg Asn Pro Ser Phe Val Leu Ser Ser Leu Gly Phe Thr Ala 8

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Val Ala Phe Val Thr Gly Ser Leu Ala Leu Trp Ala Pro Ala Phe Leu 100 105 85 9

Ę λg Ser Arg Val Val Leu Gly Glu Thr Pro Pro Cys Leu Pro Gly 115 120 125

Leu Thr Gly Val Leu Gly Val Gly Leu Gly Val Glu Ile Ser Arg Arg 145 150 150 Ser Cys Ser Ser Ser Asp Ser Leu Ile Phe Gly Leu Ile Thr Cys 130 140

5

Asp

Leu Arg His Ser Asn Pro Arg Ala Asp Pro Leu Val Cys Ala Thr Gly
165 170 175

5

Gly Ser Leu Leu Gly Ser Ala Pro Phe Leu Phe Leu Ser Leu Ala Cys Ala Arg 180 Tyr Ile Phe Ile Phe Ile Gly Glu Thr Leu 200

20

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Leu Ser Met Asn Trp Ala Ile Val Ala Asp Ile Leu Tyr Val Val 210 220 Ile Val Ala Thr 195

Ile Pro Thr . 225 ₽rg Arg Ser Thr Ala Glu Ala Phe Gln Ile Val Leu Ser 230 235

His Leu Leu Gly Asp Ala Gly Ser Pro Tyr Leu Ile Gly Leu Ile Ser 245 250 255

30

₽ ğ Leu Arg Arg Asn Trp Pro Pro Ser Phe Leu Ser Glu Phe Arg 260 265

Ala Leu Gln Phe Ser Leu Met Leu Cys Ala Phe Val Gly Ala Leu Gly 275 $280\,$

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8 Gly Ala Leu Pro Gly His Arg His Leu His Xaa 290 295

(2) INFORMATION FOR SEQ ID NO: 390:

45

Ξ SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 49 amino acids

Met Gly Pro Gln Gly Trp Val Arg Pro Leu Lys Thr Ala Pro Lys Leu 1 5 X. SEQUENCE DESCRIPTION: SEQ ID NO: 390:

50

SS Gly Glu Ala Ile Arg Leu Ile Leu Phe Leu Asn Phe Val Lys Gln Cys 20 25

Ile Ala Ser Val Asn Leu Cys Ile Leu Arg Leu Asn Ile Thr Pro Leu 35 † 40

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INFORMATION FOR SEQ ID NO: 391: (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 61 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

5

Met Tyr Val Asn Tyr Gly Thr Arg Asn Tyr Ser Thr Glu Gly Pro Ala 1 5 X. SEQUENCE DESCRIPTION: SEQ ID NO: 391:

5

Phe Ala Leu Leu Val Leu Leu Phe Val Cys Phe Cys Gly Leu Ser Tyr Val Val Ile 35 $40\,$ Asp Gln Ala Lys Leu Ser Leu Leu Val Trp Val Leu Cys 20 25 30

Ala Gln Val Pro Val Gly Leu Leu Cys Ile Thr Glu Xaa 50 55 60

25

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(2) INFORMATION FOR SEQ ID NO: 392:

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Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 392: SEQUENCE CHARACTERISTICS (A) LENGTH: 79 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid

Met Leu Trp Phe Ala Asn Phe Phe Thr Tyr Leu Phe Leu Ser Gln Ser 1 10

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Val Ala Phe Val His Ile Ser His Ile Gly Val Arg Gln Val Asn Thr 25

8

Asn Cys Tyr Phe Ser Arg Lys Ser Tyr Cys Tyr Gly Ile Leu Asn Pro 45

3

Lys Lys Lys Ile Pro Ala Gly Arg Xaa Leu Phe Pro Phe Gly 65 70

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INFORMATION FOR SEQ ID NO: 393:

9 SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 36 amino acids

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 393:

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Arg Arg Ser Glu Ser Cys Ser Met Leu Pro Trp Xaa Ala Gln 115 Pro Ala Phe Ser Arg Thr Ser Pro Trp Arg Ser Pro Lys Asn Phe 150 Arg Arg Leu Tyr Pro Pro Cys Thr Thr Ser Gly Cys Ala Ala Arg Trp 175 Met Ala Gin Ser Arg Asp Gly Gly Asn Pro Phe Ala Glu Pro Ser Glu Gln Pro Ser Arg Lys Leu Ser Pro Thr Glu Pro 70 75 80 Lys Asn Tyr Gly Ser Tyr Ser Thr Gln Ala Ser Ala Ala Ala Ala Thr 95 Ala Glu Leu Leu Lys Lys Gln Glu Glu Leu Asn Arg Lys Ala Glu Glu 100 Leu Asp Arg Thr Ile Gly Pro Leu Tyr Leu Leu Phe Val Gln Phe 130 Leu Asp Asn Pro Phe Gln Asp Pro Ala Val Ile Gln His Arg Pro Ser 20 Arg Gln Tyr Ala Thr Leu Asp Val Tyr Asn Pro Phe Glu Thr Arg Glu 35 Leu Leu Ser Leu Ser Tyr Ser Thr Leu Ser Gly Val 25 Leu Pro Pro Pro (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 394: Pro 60 Met Pro Gly Ala Phe Ser Glu Thr Val Ile Asn Asp 1 5 10 Pro Pro Pro Ala Tyr Glu Pro Pro Ala Pro Ala (A) LENGTH: 180 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 394: Pro Ala Glu Leu Ser Ala Pro Ser Leu Xaa Phe Ser Xaa 180 3 2 Tyr Arg Asn Ala 35 Phe Leu Val 3 ပ္ပ Leu Asp Ser 145 9 2 S 55 35 6 5 S 2 8 22 ഉ 2

Met Fro Thr Pro Cys Thr Ser Leu Pro Ser Cys Cys Gln His Arg Ser $1 \\ 1 \\ 1$ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 395: (A) LENGTH: 21 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear lle Thr Met Thr Leu 2

(2) INFORMATION FOR SEQ ID NO: 396:

15

Ser Lys His Pro Ile Gin Met Ser Leu Cys Met Cys Val Asn Ile $20 \ 25$ Pro Leu Phe Ile Pro Leu Ile Phe Phe Leu Ser Leu Leu His Cys 10 SEQUENCE DESCRIPTION: SEQ ID NO: 396: (A) LENGTH: 60 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear (X Ê Met g Ser 20 52

Leu Val Trp Ser Pro Val Arg Trp lle Phe Cly Ser Lys Cly Leu 35 45 Val His Leu Gln Ser Ser Gln Arg Pro Ser 55 60 Phe 8

(A) LENGTH: 152 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 397: 6

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 397:

Ile Ile Val Leu Val Ile Val Val Ile Cys Leu Met Leu Tyr Ala Leu $20\ \ 25$ Leu Trp Glu Ala Gly Asn Leu Thr Asp Leu Pro Asn Leu Arg Ile Gly 35 Met Ala Gly Pro Arg Pro Xaa Trp Arg Asp Gln Leu Leu Phe Met Ser 1 45 S

His Gln Phe Pro Glu Leu Glu Ala Leu Gly Val Pro Arg Val Gly Leu 65 70 80 Phe Tyr Asn Phe Cys Leu Trp Asn Glu Asp Thr Ser Thr Leu Gln Cys 50 60 55

Gly Leu Ala Arg Leu Gly Val Tyr Gly Ser Leu Val Leu Thr Leu Phe \$95\$

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(1) SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 395:

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Thr Asp Asp Ala Leu Gln Glu Leu Val Glu Leu Ile Tyr Gln Gln Ala

8 Glu Thr Glu Ile Glu Gln Phe Ala Glu Thr Leu Asn Gly Cys Val Thr 180

55 Tyr Val Gln Asp Phe Leu Asn His Leu Thr Glu Gln Pro Gly Ser Phe 165 170 175

The Glu Ser Tyr Glu Asp Gly Cys Glu Asp Tyr Pro The Leu Ser Glu 145 150

Ser Val Asn Ala Pro Glu Phe Tyr Pro Ser Gly Tyr Ser Ser Ser Tyr 130 140

Met Ala Lys Pro Gin Val Val Ala Pro Val Leu Met Sex Lys Leu 115 120 125

Pro Ser

Ser Gln Asp Lys Ile Pro Gln Gln Asn Ser Glu Ser Ala 100 105 110

Pro Ser Arg Pro Gly Ala Leu Pro Glu Gln Thr Arg Pro Leu Arg Ala 95

6

35

Thr Pro Pro Pro Gly Ala Gln Cys Glu Val Pro Ala Ser Pro Gln Arg 65 70 75 80

Lys Ala Pro Gly Phe Leu Gln Pro Xma Pro Leu Arg Gln Pro Arg Thr 50 55

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Xaa Gly Xaa Xaa Pro Ala Glu Arg Xaa Arg His Gln Pro Pro Gln Pro 35 40 45

3

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5 S Gly Gly Leu Gly Leu Phe Leu Ser Tyr Val Trp Asn Gly Ser Xaa Ser 130 140 Trp Arg Leu Ala Val Gly Phe Leu Ala Val Ser Ser Val Leu Leu Ala 115 120 125 Ala Pro Gl
n Pro Leu Leu Leu Ala Gl
n Cys Asn Xaa Asp Glu Arg Ala 105 $^{\circ}$ 110

Pro Ser Arg Gly Leu 145 1 Gly Phe Xaa 150

3 INFORMATION FOR SEQ ID NO: 398:

5

ž E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 398: (A) LENGTH: 480 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

20

Gln Val Thr Arg Ala Asp Ile Leu Gln Val Gly Leu Arg Glu Leu Leu 290 300

Phe Leu Gly Glu Leu Tyr Leu Asn Leu Glu Ile Lys Gly Thx Asn Gly 275 280

Asn Ala Leu : 305

Phe Ser Asn Pro Met Asp Asp Asn Leu Ile Cys Ala Val 310 315

2

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Leu Leu Gin Arg Cys Arg Thr Glu Tyr Glu Val Lys Asp Gin Ala 255 245

ala Lys Gly Asp Glu Val Thr Arg Lys Arg Phe His Ala Phe Val Leu 265 265

Leu Ser His His Leu Thr Ile Ser Pro Gln Ser Gly Asn Phe Ary Gln 225 230 230

Thr Ser Ile Pro Asn Phe Ser Tyr Met Gly Ala Arg Leu Cys Asn Tyr 210 215 220

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200

205

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Met Ser Asp Gly Phe Asp Arg Ala Pro Gly Ala Gly Arg Gly Arg Xaa 1 10 15

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Arg Gly Leu Gly Arg Gly Gly Gly Gly Pro Xaa Gly Gly Gly Phe Pro 20 25 30

25 Lys Leu Leu Lys Leu Thr Gly Ser Val Leu Glu Asp Ala Trp Lys Glu 325 330 335

Lys Gly Lys Met Asp Met Glu Glu Ile Ile Gln Axg Ile Glu Asn Val 340°

Val Leu Asp Ala Asn Cys Ser Arg Asp Val Lys Gln Met Leu Leu Lys 355 360

30

Leu Val Glu Leu Arg Ser Ser Asn Trp Gly Arg Val His Ala Thr Ser 370 375 380

Thr Tyr Arg Glu Ala Thr Pro Glu Asn Asp Pro Asn Tyr Phe Met 385 390 400

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Glu Pro Thr Phe Tyr Thr Ser Asp Gly Val Pro Phe Thr Ala Ala Asp 415

6

Pro Asp Tyr

Gln Glu Lys Tyr Gln Glu Leu Clu Arg Glu Asp Pho 420 425

Pro Asp Tyr Glu Glu Asn Gly Thr Asp Leu Ser Gly Ala Gly Asp 435

3

Phe 15 Y Leu Asp Asp Ile Asp Asp Glu Met Asp Pro Glu Ile Glu Glu 455

Ala Tyr Glu Lys Phe Cys Leu Glu Ser Glu Arg Lys Arg Lys Gln Xaa 465 470 475 Pro

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INFORMATION FOR SEQ ID NO: 399:

592

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lle fhr Leu Met Gly Pro Gln Trp Trp Leu Lys Thr Val Ile Glu Gln 290 296

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PCT/US98/04493 Leu His Ser Tyr Leu Leu Gly Asp Gln Glu Glu Asn Glu Asn Ser Ala Asn Gln 85 95 Ser Leu Ile Cys Leu Thr Leu Pro Val Phe Ala Gly Arg Trp Leu Met 175 175 Cys Gly Leu Tyr Val Cys Trp Leu Thr lle Arg Ala Val Thr Val Met 195 Val Ala Trp Met Pro Gln Gly Arg Arg Val 11e Phe Gln Lys Val Lys 210 Gly Pro Val Gly Phe Gln Xaa fyr Arg Arg Pro Leu Asn Phe Pro Leu 130 Arg lie Phe Leu Leu Ile Val Phe Met Cys lle Thr Leu Leu Ile Ala 145 Ser Fhe Try Thr Gly Thr Ala Lys Ile Hie Glu Leu Tyr Thr Ala Ala 185 Met Glu Pro Lys Thr Ile Thr Asp Ala Leu Ala Ser Ser Ile Ile Lys Leu Pro Asn Phe Leu Pro Tyr Asn Val Met Leu Tyr Ser Asp 20 Pro Val Ser Glu Leu Ser Leu Glu Leu Leu Leu Gln Val Val 35 pro Ala Leu Leu Glu Glu Gly His Thr Arg Gln Trp Leu Lys Gly 50 60 Gin Val Asn Asn Gin His Ala Arg Asn Asn Asn Ala Ile Pro Val 100 Val Gly Glu Gly Leu His Ala Ala His Gln Ala Ile Leu Gln Gln Gly 115 Leu Val Arg Ala Trp Thr Val Thr Ala Gly Tyr Leu Leu Asp 65 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 399: 593 (A) LENGTH: 423 amino acids SEQUENCE CHARACTERISTICS TYPE: amino acid (B) TYPE: amino acid (D) TOPOLOGY: linear 3 Ser Val WO 98/39448 Ę 35 25 승 5 2 2 ഉ 2

Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Aan Pro Ser Gly Pro 50Leu Arg Leu Asp Ile Ile Asn Ser Leu Val Thr Thr Val Phe Met 5 Val Gly Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 35 46 Leu Val Asn Tyr Glu Arg Lys Ser Gly Lys Oln Gly Ser Ser Pro Pro 405 415 Vel Tyr Ala Asn Gly Ile Arg Asn Ile Asp Leu His Tyr Ile Val Arg 310 310 Leu Leu Leu Ser Leu Cys Val 330 Val Val Vel Leu Met Ala Ile Leu Ser Phe Gln Val Arg Gln Phe Lys 370 Arg Leu Tyr Glu His Ile Lys Asn Asp Lys Tyr Leu Val Gly Gln Arg 185 Leu Thr Pro Tyr Val Ile Ala Ser Gly Val Val Pro Leu Leu Gly Val Thr Ala 345 Leu Leu Met Leu ile Val Ser Val Leu Ala Leu ile Pro Glu Thr Thr Thr 25 20 20 Tyr Gln Lys Lys Pro Val His Glu Lys Lys Glu Val Leu Xaa 65 Glu Met Gln Asn Leu Val His Arg Arg Ile Tyr Pro Phe 360 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 400: (A) LENGTH: 78 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: Lys Leu Ala Ala Pro Val Ile Ser Val 325 INFORMATION FOR SEQ ID NO: 400: Pro Pro Gln Ser Ser Gln Glu 420 <u>2</u> Met S 45 9 35 **4** 8 9 2 22

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 74 amino acids
(B) TYPE: amino acid
(D) TOPOLGY: linear

8

Trp Gln Asp Trp Ala Leu Gly Val Leu His Ala Lys Ile Ile Ala Ala 275

8

Ala Gly Val Val Pro Leu Leu Leu Gly Leu Leu Phe Glu Leu Val Ile 250

Glu Trp Ser Leu Met Ile Met Lys Thr Leu Ile Val Ala Val Leu Leu 240 235

S

Val Ala Pro Leu Arg Val Pro Leu Asp Gln Thr Pro Leu Phe Tyr Pro 260.

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(2) INFORMATION FOR SEQ ID NO: 401:

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 401:

S His Cys Trp Gly Leu Pro Leu His Val Ala Pro Leu Cys Arg Gly His 20 25 Met Leu Lys Gln Val Met Phe Val Phe Ser Gly Met Gly Pro Arg Ser 1 10

5 Gln Ala Asp Ser Ser His Leu Leu Pro Leu Lys His Gln Gly Ala Trp 35 40

5

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(2) INFORMATION FOR SEQ ID NO: 404:

(1) SEQUENCE CHARACTERISTICS:

(A) LENOTH: 92 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid

Asn Arg Asn Leu Ala Asn Gln Arg His Phe Phe Cys Pro Ser Ile Phe 50 55

His The Cys Pro Val Leu Phe Phe Xaa 70

2

S

(2) INFORMATION FOR SEQ ID NO: 402:

20

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 402: (B) TYPE: amino acid
(D) TOPOLOGY: linear

25

Ala Arg Thr Ile Leu Val Leu Tyr Leu Ser Leu Gln Arg Leu Glu Asn 1 15

Leu Ala Tyr His 20

(2) INFORMATION FOR SEQ ID NO: 403:

35

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 87 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

6

Met Pro Leu Pro Ser Val Pro Ile Leu Gly Ile Phe Ser Phe Leu Ile 1 5 (X SEQUENCE DESCRIPTION: SEQ ID NO: 403:

3 Pro. Ser Ser Gln Gly Val Ser Tyr Thr Lys Leu Pro Ile Ser Ser Pro 20 25 30

8

Pro Val Gln Ile His Thr Gly Phe Ala Arg Val Gly Ser Tyr Met Gln 50 55

55 Met Pro Leu Val His Leu Cys Leu Leu Gln Thr Ser Leu Met Lys Aøn 80 75

Ser Gly Val Gln Gln Gly Ser

8

Gln Tyr Ser Pro Phe Val Asn Asp His Phe Ser Phe Leu Asn Pro Phe 35

30

INFORMATION FOR SEQ ID NO: 405:

(1) SEQUENCE CHARACTERISTICS: (A) LENCTH: 21 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 405:

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8 Met Ala Cys Ser Cys Leu Met Ile Gln Ser Phe Ser Thr Ser Ala Leu 1 15

Val Leu Phe Tyr Gly 20

45

(1) SEQUENCE CHARACTERISTICS:

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(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 174 amino acids

Met Glu Glu Gly Gly Asn Leu Gly Gly Leu Ile Lys Met Val His Leu 1 15

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Gln Thr Lys Ser Ile Val Glu Lys Ile Pro Ser Lys 85

8

Leu Leu Ile Leu Ala Glu Leu Leu Ile Pho Ser Val Pro Ile Val Tyr 50 55

Glu Lys Tyr Lys Thr Gln Ile Asp His Tyr Val Gly Ile Ala Arg Asp 65 70 75

15

 $\mbox{Arg Leù Phe Leu Val Glu Asp Leu Val Asp Ser Leu Lys Leu Ala Val <math display="inline">\mbox{20}$

Phe Met Trp Leu Met Thr Tyr Val Gly Ala Val Phe Asn Gly Ile Thr 35

Met Asn Ala Ala Met Val His Ile Asn Arg Ala Leu Lys Leu Ile Ile 1 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 404:

2

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 406:

S

Leu Val Leu Ser Gly Ala Trp Gly Met Gln Met Trp Val Thr Phe Val 20 25 10

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PCT/US98/04493

Ser Gly Phe Pro Ala Phe Pro Lys Pro Ser Pro Thr Leu Arg Thr 15 45

Ser Ala Glu Gln Thr Leu Pro Leu Leu Pro His Leu His Gly Leu 50 60

Leu His Gln Pro Leu His Leu Gly Phe Thr Ala Cys Leu Gly Ser $$75\$ 8 % 2

Ala His Ile Leu Gly Gly Gln Pro Ala Leu Pro Ala Val Pro Glu Pro 95

Tyr Ala Gly His Cys Gln Arg Pro Leu Ala Gly Thr Pro His His Ser 100

2

Gly Arg Tyr Gln Ala Ala Asn Arg Phe Pro Ile Leu Asn Ala Xaa Cys 130 Cys His Val Gly Pro Ala Asn Arg Gly Arg Arg Ser Glu Ala Trp Val 115 ឧ

Ser Ala 160 Glu Arg Arg Thr Pro Ser Thr Val Leu Ser Ala Arg Ile Ser 145

Thr Met Gly Cys Pro Leu Phe Ala Ile Trp Ala Ala Ser Xaa 170

23

(2) INFORMATION FOR SEQ ID NO: 407:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 407: (D) TOPOLOGY: linear

Glu Gln Asn Ser Ala Thr Val Glu Pro Ser Ser His Glu Ile Leu His 30 Met Ala Phe Ile Leu Leu Phe Tyz Cys Leu Met Thz Phe Leu Ser Leu 1 10 15 6

Leu Gln Asn Cys Phe Glu Leu Leu Arg Thr Ser Thr Ser Gln Cys 35 <u>Leu</u> 45

Glu Gly Ile Pro Cys Gln Arg fyr Gln Asn Gly Leu His Ile Xaa 50 뀹 S

(2) INFORMATION FOR SEQ ID NO: 408:

55

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 280 amino acids
(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 408: (D) TOPOLOGY: linear

Met Glu Ala Val Val Asn Leu Tyr Gln Glu Val Met Lys His Ala Asp Pro Arg Ile Gln Gly Tyr Pro Leu Met Gly Ser Pro Leu Leu Met Thr $20 \ 25 \ 30$ Ser Ile Leu Leu Thr Tyr Val Tyr Phe Val Leu Ser Leu Gly Pro Arg 35 9

Ile Met Ala Asn Arg Lys Pro Phe Gln Leu Arg Gly Phe Met Ile Val $50\,$

2

Tyr Asn Phe Ser Leu Val Ala Leu Ser Leu Tyr Ile Val Tyr Glu Phe 65 Leu Met Ser Gly Trp Leu Ser Thr Tyr Thr Trp Arg Cys Asp Pro Val 85 20

Asp Tyr Ser Asn Ser Pro Glu Ala Leu Arg Met Val Arg Val Ala Trp 110

Leu Phe Leu Phe Ser Lys Phe Ile Glu Leu Met Asp Thr Val Ile Phe. 125 ile Leu Arg Lys Lys Asp Gly Gin Val Thr Phe Leu His Val Phe His 130

23

His Ser Val Leu Pro frp Ser Trp frp Gly Val Lys Ile Ala Pro-145

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Gly Gly Met Gly Ser Phe His Ala Met Ile Asn Ser Ser Val His Val. 175 35 Ile Met Tyr Leu Tyr Tyr Gly Leu Ser Ala Phe Gly Pro Val Ala Glni 180 Pro Tyr Leu Irp Irp Lys Lys His Met Thr Ala Ile Gln Leu Ile Gln 195, 195,

6

Phe Val Leu Val Ser Leu His Ile Ser Gln Tyr Tyr Phe Met Ser Ser 210

Cys Asn Tyr Gin Tyr Pro Val 11e Ile His Leu 11e Trp Met Tyr Gly, 225 $\,$ S 45

Lys Gly Lys Arg Leu Pro Arg Ala Leu Gln Gln Asn Gly Ala Pro Gly 260 260 Thr lle Phe Net Leu Phe Ser Aan Phe Trp Tyr His Ser Tyr Thr 255 255

Ile Ala Lys Val Lys Ala Asn Xaa 280

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60 . (2) INFORMATION FOR SEQ ID NO: 409:

PCT/US98/04493

WO 98/39448

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 284 amino acids

(XX SEQUENCE DESCRIPTION: SEQ ID NO: 409: (B) TYPE: amino acid (D) TOPOLOGY: linear

S

Met Xaa Leu ' Trp Pro Gln Thr Cys Ser Gly Lys Phe Asp Gly Thr Leu 5

Ala Phe Ser Ile His Xaa Leu Ala Val Ile Leu Gly Asp 20 25 Gln Leu Thr 30

5

15 Glu Val Arg Ile Gly Val Leu Lys His Leu 50 55 Ala Ala Asp Leu Val Pro Ile Phe Asn Gly Phe Leu Lys Asp Leu Asp 45His Asp Phe Leu Lys Leu 60

20 Leu His Ile Asp Lys Arg Arg Glu Tyr Leu Tyr Gln Leu Gln Glu Phe 65 70

Leu Val Thr Asp Asn Ser Arg Asn Trp 85 Arg Phe Arg Ala Glu Leu Ala 90 95

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Glu Leu Lys Gln Arg Phe Ser Val Phe Gly Glu Ile Glu Glu Cys Thr $90\,$

Arg Arg Val Val Phe Ile Gly Lys Ile Pro Gly Arg Met Thr Arg Ser 65 70 75

20

His Tyr Gln Arg Gln Arg Val Leu Gln Lys Glu Arg Ala Ile Glu Glu 50 60

Gly XAA Ser Asp Arg Arg Arg Tyr Ser Ser Tyr Arg Ser His Asp 35

25 Glu Gln Leu Ile Leu Leu Leu Glu Leu Tyr Ser Pro Arg Asp Val Tyr 100 105

30 Asp Tyr Leu Arg Pro Ile Ala Leu Asn Leu Cys Ala Asp Lys Val Ser 115 120

Ser Val Arg Trp Ile Ser Tyr Lys Leu Val Ser Glu Met Val Lys Lys 130 135

Leu His Ala Ala Thr 145 Pro Pro Thr Phe Gly Val Asp Leu Ile Asn Glu 150 155 160

35

Leu Val Glu Asn Phe Gly Arg Cys Pro Lys Trp Ser Gly Arg Gln Ala 175

Phe Val Phe Val Cys Cln Thr Val Ile Glu Asp Asp Cys Leu 180 185 Pro Met

6

Asp Gln Phe Ale Val His Leu Met Pro His Leu Leu Thr Leu Ale Asn 195 200 205

25

βşρ Arg Val Pro Asn Val Arg Val Leu Leu Ala Lys Thr Leu Arg Gln 210 220

Thr Leu Leu Glu Lys Asp Tyr Phe Leu Ala Ser Ala Ser Cys His Gln 240 225

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Glu Ala Val Glu Gln Thr Ile Met Ala Leu Gln Met Asp 250 Asp Val Lys Tyr Phe Ala Ser Ile His 265 Pro Ala Ser 쿭 : Lys Ile Ser 270 Asp Ser 255

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5 U Asp Ala Met \$er Thr Ala Ser Ser Thr Tyr Xaa 275

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'n (2) INFORMATION FOR SEQ ID NO: 410: Ξ

X SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: 410: (A) LENGTH: 187 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Phe Val Phe Leu

5 Met Leu Phe Leu Phe Phe Val Ile Ile Phe Leu 1 5 10 Leu Ile Ile Gln Phe Ser Lys Pro Leu Thr Asn Pro His Pro Pro Ala 20 25 30 15 Ile

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4 Gln Ala Asp Glu Gln Pro Phe Asp Leu Cys Phe Gly Gly Arg Arg Xaa

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Tyr Ala Glu Glu Ala Phe Ala Ala Ile Glu Ser Gly His Lys Leu Arg 115 126

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Ile His Phe

Arg Val Gln Gly Asp Asn Tyr Gly Phe Val Thr Tyr Arg 100 105

Xaa Cys Lys Arg Ser Tyr Ser Asp Leu Asp 145 Ser Asn Arg Glu Asp Phe 155

gg Gg Pro Ala Pro Val Lys Ser Lys Phe Asp Ser Leu Asp Phe Asp Thr 165 170

3 Leu Leu Lys Gln Ala Gln Lys Asn Leu Arg Arg 180 185

50 (2) INFORMATION FOR SEQ ID NO: 411:

Ξ SEQUENCE CHARACTERISTICS acids

£ SEQUENCE DESCRIPTION: SEQ ID NO: 411: (A) LENGTH: 237 amino a (B) TYPE: amino acid (D) TOPOLOGY: linear

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Met Lys Leu Pro Gly Lys Phe Arg Arg Ala His Gln Gly Asn Leu Glu 1 15

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Ala Pro Xaa 50 Glu Gln Val Phe Leu Val Ile Leu Ala His Val Val Arg Arg Cys Ala $20 \ \ 30$ Met Leu Ile Ile Ser Leu Arg Pro Gln Phe Pro Ser Leu Ile Val Gln 1 5 10 Ser Asp Gly Ile Leu Gln Phe Glu Ser Ser Leu Leu Lys Met Arg Arg 35 (2) INFORMATION FOR SEQ ID NO: 415: Ξ (X SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 415: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 32 amino acids

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25 20 Leu Glu Cys Ser Val Leu Phe Leu Pro Ile Ser Leu Asn Leu Leu Leu 20 30

30 (2) INFORMATION FOR SEQ ID NO: 416:

Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 416: SEQUENCE CHARACTERISTICS (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 163 amino acids

35

6 Met Val Lys Val Cys Asn Asp Ser Asp Arg Trp Ser Leu Ile Ser Leu 1 15 Ser Asn Asn Ser Gly Lys Asn Val Glu Leu Lys Phe Val Asp Ser Leu 20 30

2 Arg Arg Gln Phe Glu Phe Ser Val Asp Ser Phe Gln Ile Lys Leu Asp 35 40 45

Ser Leu Leu Phe Tyr Glu Cys Ser Glu Asn 50 55 Pro Met Thr Glu Thr 60

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55 Phe His Pro Ala Phe Asp His Leu Cys Asn Lys Ile Ile Ala Thr Arg Asn Pro Glu 85 90 95

Glu Ile Arg Gly Gly Gly Leu Leu Lys Tyr Cys Asn Leu Leu Val Arg 100 105

ප Gly Phe Arg Pro Ala Ser Asp Glu Ile Lys Thr Leu Gln Arg Tyr Met

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ş Ser Arg Phe Phe Ile Asp Phe Ser Asp Ile Gly Glu Gln Gln Arg 130 140 15 120 125

5 Gln Val

Lys Leu Glu : 145

Ser Tyr Leu Gln Asn His Phe Val Gly Ile Gly Arg Pro 150 155

(2) INFORMATION FOR SEQ ID NO: 417:

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(i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 417: (A) LENGTH: 174 amino acids
(B) TYPE: amino acid
(D) TOPOLOCY: linear

25 20 Met Ala Pro Lys Gly Lys Val Gly Thr Arg Gly Lys Lys Gln Ile Phe 1 15

30 Ala Ser Phe Trp Ala Trp Leu Ala Leu Gly Phe Ser Leu Ala Val Tyr S0 $$ 55 $$ 60 $$. Ala Asn Ala Ile Tyr Cys Leu Val Thr Leu Val Phe Phe Tyr Ser Ser 35 40 45 Glu Glu Asn Arg Glu Thr Leu Lys Phe Tyr Leu Arg Ile Ile Leu Gly $25\ \ 30$

Glu Asp Gly Ala Leu Met Asp Gly Gly Met Asp Leu Asn Met Glu Gln
95 Gly Ala Ser Tyr His Ser Met Ser Ser Met Ala Arg Ala Ala Phe Ser 65 70 75 80

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Gly Met Ala Glu His Leu Lys Asp Val Ile Leu Leu Thr Ala Ile Val 100' 105 110

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Ala Pro Gly Arg Ala Leu Tyr Leu Leu Trp Val Asn Val Leu Gly Pro 130 Gln Val Leu Ser Cys Phe Ser Leu Tyr Val Trp Ser Phe Trp Leu Leu 115

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Trp Phe Thr Ala Asp Ser Gly Thr Pro Ala Pro Glu His Asn Glu Lys 145 150 150 Arg Gln Arg Arg Gln Glu Arg Arg Gln Met Lys Arg Leu Xaa 165 170

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INFORMATION FOR SEQ ID NO:

2)

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(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 50 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 418: (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Glu Leu Pro Lys Gly Leu Gln Gly Val Gly Pro Val Ala Met Met 1 15 1

Leu Pro Val Leu Cys Thr Gln Ala Leu Arg 25 ž Arg Pro Phe Tyr Leu 20

Ser Gln Gly Lys Ser Pro Leu Leu Trp Lys Arg Thr Cys Cys Leu 35 40 gju

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Ala Xaa 50

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(2) INFORMATION FOR SEQ ID NO: 419: ಣ 3

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 120 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 419:

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Leu Tyr Phe Thr Leu Val Val Gly Glu Gly Glu Pro Gly Glu Aan Lys 20

Thr Gly Lys Lys Ile Ile Phe Cys Ser 40

Phe Glu

Thr Ile Pro Phe 35

Ser Asn Val Pro Ser His Lys Gly Pro Val 55 60 Val Lys Met Val Glu Asn 35

Leu Gly 80 Pro Leu Arg Ser Glu Gln Trp Glu Leu Lys Ile Ser Glu Thr 65 75 4

Phe Leu Leu Ile Gly Arg Cys Ser Ser Gly Xaa 90 Leu Cys Phe Cys Trp Asp Val Leu Cys Cys Met Tyr Ala Tyr 100 Glu Gly Lys Ile Gly 85 먑 45

12 E Arg Ser Leu Leu Ser 115 ABD Met

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INFORMATION FOR SEQ ID NO: 420 3

(A) LENGTH: 159 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

22

SEQUENCE DESCRIPTION: SEQ ID NO: 420: Z

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Lys Ala Val Pro Val Ala Pro Glu Val Thr Gln Gln Thr 11e Glu Leu Lys Glu Glu Cys Lys 50 60 Leu Arg Val Leu 45 Asp Phe Val Asp Lys Ile Gly Gln Phe Gln Lys Ile Val Gly Gly Leu 70 75 80 Met Asn Val Gly Val Ala His Ser Glu Val Asn Pro Asn Thr Arg Val Met Aen Ser Arg Gly Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu 25 Ser Ser Arg Lys Phe Phe Thr 11e Ser Pro 11e 11e Leu Tyr Phe Leu 110 Met Thr His Leu Leu Leu Thr Ala Thr Val Thr Pro Ser Glu Gln Asn Ser Ser Arg Glu Pro Gly Trp Glu Thr Ala Met Ala Lys Asp Ile Leu 20 lle Glu Leu Val Asp Gln Leu Ala Lys Glu Ala Glu Asn Glu Lys Met 95 Lys Ala Ile Gly Ala Arg Agn Leu Leu Lys Ser Ile Ala Lys Gln Arg 110 Glu Ala Gln Gln Gln Gln Leu Gln Ala Leu Ile Ala Glu Lys Lys Met 115 Gln Leu Glu Arg Tyr Arg Val Glu Tyr Glu Ala Leu Cys Lys Val Glu 130 Trp Thr Leu Thr Asn Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu Arg Leu Leu Thr His Trp Glu Gln Leu Asp Tyr Gly Val Gln Phe Thr 95 Ala Glu Gln Aan Glu Phe Ile Asp Gln Phe Ile Phe Gln Lys Xaa 145 Thr Pro Asp Gln Gly SEQUENCE DESCRIPTION: SEQ ID NO: 421: Gly Glu Ala Gly Leu His Phe Asp Glu Leu Asn Lys Ser Ile Pro Phe Phe Ser (A) LENGTH: 154 amino acids Thr Pro Phe Glu 70 SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 421: (D) TOPOLOGY: linear Leu His Ile Val Leu Leu His Ala Val Lys Gly (xi Asp 8 2 2 20 පි 15 23 8 35 송 5 55

Ala Ser Leu Leu Ser Val Leu Ile Pro Lys Met Pro Gln Leu His Gly 130 140 Val Arg Ile Phe Gly Ile Asn Lys Tyr Xaa 145 Ala Ser Phe Tyr Thr Lys Tyr Asp Pro Thr His Phe Ile Leu Asn Thr 115 120 Cys Gly Phe Arg Ser Val Asn Pro 130 135 Val Ile Ala Val Gly Ile Phe Leu Phe Leu Ile Ala Leu Val Gly Leu 50 55 Trp Gly Ile Gly Phe Gly Leu Ile Ser Ser Leu 35 40 Met Val Cys Gly Gly Phe Ala Cys Ser Lys Asn Cys Leu Cys Ala Leu 1 15 (2) INFORMATION FOR SEQ ID NO: 422: Gln Lys Asp Pro Arg Ala Asn Pro Ser Ala Phe Leu 195 Tyz Ala Gly Glu Val Leu Arg Phe Val Gly Gly Ile Gly Leu Phe Phe 175 $$170\,$ Val Lys Ser Asp His Ser Cys Ser Pro Cys Ala Pro Ile Ile Gly Glu 145 150 155 Asn Asn Thr Ala Ser Ala Arg Asn 115 Ile Leu Leu Leu Val Phe Ile Val Gin Phe Ser Val Ser Cys Ala Cys
95 Ile Gly Ala Val 65 Asn Leu Leu Leu Ala Leu Phe Thr Glu Ile Leu Gly Val Trp Leu Thr Tyr Arg Tyr Arg Asn 180 185 E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 422: Asn Gin Glu Gin Gly Gin Leu Leu Glu Val Gly Trp 100 105 110 Tyr Thr Leu Val Ser Leu Leu Leu Ile Gly Ile Ala Ala 20 25 30 SEQUENCE CHARACTERISTICS: (A) LENGTH: 204 amino acids
(B) TYPE: amino acid Lys His His Gln Val Leu 70 Asp Ile Gln Arg Asn Leu Asn Cys 125 λgn Asp Thr Cys Leu Ala Ser Cys 140 Leu Phe Phe Tyr Met Ile 75 80 ξ val val Gly val 45

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(2) INFORMATION FOR SEQ ID NO: 423:

Ē (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 423: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 67 amino acids

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5 Met Leu Gln Ser Ile Ile Lys Asn Ile Trp Ile Pro Met Lys Pro Tyr
1 10 15

5 Tyr Thr Lys Val Tyr Gln Glu Ile Trp Ile Gly Met Gly Leu Met Gly
20 25 10

Phe Ile Val Tyr Lys Ile Arg Ala Ala Asp Lys Arg Ser Lys Ala Leu 45Lys Ala Ser Ala Pro Ala Pro Gly His His Asn Gin Ile Tyr Leu Glu 50 60

Tyr Met Xaa 65

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23

(2) INFORMATION FOR SEQ ID NO: 424:

30 E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 424: SEQUENCE CHARACTERISTICS: (A) LENGTH: 25 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

ઝ Met Leu Gly Val Ser Leu Phe Leu Leu Val Val Leu Tyr His Tyr Val 1 15

Ala Val Asn Asn Pro Lys Lys Gln Glu 20 25

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(2) INFORMATION FOR SEQ ID NO: 425:

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£ $\widetilde{\mathbf{x}}$ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 425: (A) LENGTH: 299 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

Met Ala Ala Xaa Glu Pro Ala Val Leu Ala Leu Pro Asn Ser Gly Ala 1 15

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55 Gly Gly Ala Gly Ala Pro Ser Gly Thr Val Pro Val Leu Phe Cys Phe
20 25 30 Ser Val Phe Ala Arg Pro Ser Ser Val Pro His Gly Ala Gly Tyr Glu
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Leu Leu Ile Gln Lys Phe Leu Ser Leu Tyr Gly Asp Gln Ile Asp Met

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His Arg Lys Phe Val Val Gln Leu Phe Ala Glu Glu Trp Gly Gln Tyr 75 Ser Glu Arg Cys Lys Val Arg 90 95 Leu Pro Lys Gly Phe Ala Val 85 Val Asp

Pro Leu Gln Ile Gln Leu Thr Thr Leu Gly Asn Leu Thr Pro 100 Ser Ser Thr Val Phe Phe Cys Asp Met Gln Glu Arg Phe Arg Pro 120 Leu Val

9

Ala Ile Lys Tyr Phe Gly Asp Ile Ile Ser Val Gly Gln Arg Leu Leu 130 2

Gln Gly Ala Arg Ile Leu Gly Ile Pro Val Ile Val Thr Glu Gln Tyr 145 20

Leu Gly Ser Thr Val Gln Glu Ile Asp Leu Thr Gly Val 165 Pro Lys Gly

Lys Leu Val Leu Pro Lys Thr Lys Phe Ser Met Val Leu Pro Glu Val 180 Glu Ala Ala Leu Ala Glu Ile Pro Gly Val Arg Ser Val Val Leu Phe 195 25

Gly Val Glu Thr His Val Cys Ile Gln Gln Thr Ala Leu Glu Leu Val 210 2

Ser Ser Arg Met Met Asp Arg Met Phe Ala Leu Glu Arg Leu Ala Xaa Xaa Gly 255 Gly Arg Gly Val Glu Val His Ile Val Ala Asp Ala Thr 225 Ser

35

The The See Glu Ala Val Leu Leu Gln Leu Val Ala Asp 260 Lys Asp His Pro Lys Phe Lys Glu Ile Gln Asn Leu Ile Lys Ala Ser 275 Ile Ile Val 6

Ser Lys Val Xaa 395 295 Ze Ze Glu Ser Gly Pro 290 Ala 45

(2) INFORMATION FOR SEQ ID NO: 426: S

(A) LENGTH: 13 amino acids SEQUENCE CHARACTERISTICS: TYPE: amino acid 3

55

Met Arg Asp Leu Gly Thr Leu Leu Ser Pro Val Cys Ser 1 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 426:

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INFORMATION POR SEQ ID NO: 427 2

427: SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 198 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

(X

Phe Gly Cys Leu Val Ala Gly Arg Leu Val Gln Thr Ala Ala Gin 15 Ser Gin Val Ala Giu Asp Lys Phe Val Phe Asp Leu Pro Asp Tyr Glu $25\ \ \, 25$ Met 2 15 Ile Asn His Val Val Val Phe Met Leu Gly Thr Ile Pro Phe Pro Glu Met Met Gly Gly Ser Val Tyr Phe Ser Tyr Pro Asp Ser Asn Gly 50 60 ξ

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Pro Val Trp Gln Leu Ceu Gly Phe Val Thr Asn Gly Lys Pro Ser Ala 65

Pro Pro Ser Val Ala Gln Ile Gly 110 lle Phe Lys Ile Ser Gly Leu Lys Ser Gly Glu Gly Ser Gln His 95 7 105 Phe Gly Ala Met Asn Ile Val Arg

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Phe Thr Gln Phe Thr Gln Lys ile Ser Val Glu Leu Leu Asp Ser Met Ala Gln Gln Thr Pro Val Gly 115 Ser Asn Ala Ala Val Ser Ser Val Asp 130

130

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Met Leu Asp Asn Phe Tyr Asn Phe Ala Ser Ser Phe Ala Val Ser Gin 145 Ala Gln Met Thr Pro Ser Pro Ser Glu Met Phe Ile Pro Ala Asn Val 175 5

Val Leu Lys Trp Tyr Glu Asn Phe Gln Arg Arg Leu Ala Gln Asn Pro 190

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Xaa Phe Trp Xaa Thr Xaa 195

(2) INFORMATION FOR SEQ ID NO: 428:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 428: (A) LENGTH: 47 amino acids (B) TYPE: amino acid (1) SEQUENCE CHARACTERISTICS 22

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Met Gly Leu Pro Leu Met Ala Leu Met Trp Ser Thr Leu Pro Ala Ser

612

မ 20 5 45 35 25 5 8 6 S 5 Ala Gly Val Asn Phe Ile Leu Ala Leu Pro Leu Leu Leu Leu Trp Lys 20Ile Lys His Trp Ile Thr Ile Ile Arg Ala Arg Phe Glu Glu Val Leu \$45\$Met Gly Glu Val Ile Leu Ala Val Cys His Pro Asp Cys Ile Thr Thr 20Met Lys Lys Val Glu Glu Lys Arg Val Asp Val Asn Ser Ala Val 1 10 (2) INFORMATION FOR SEQ ID NO: 429: Asn Arg Gly Gly Val Gly Arg Ser Val Met Ser Ala Val Glu Xaa 35 40 Glu Glu Met Thr Arg Lys Gln Pro Asp Val Asp Arg Val Thr Lys Thr 115 120 125 Asn Ile Asp Arg Val Lys Ala Leu Ile Ala Glu His Gln Thr Phe Met $100\,$ Trp Ala Glu Thr Thr Leu Ile Gln Arg Asp Gln Glu Pro Ile Pro Gln 95 Leu Val Ala Asn Ala Glu Leu Leu Glu Glu Leu Leu Ala Trp Ile Gln 65 70 75 Ser Arg Ser Gly Gly Arg Lys Ser Leu Ser Gln Pro Thr Pro Pro Pro 145 Tyr Lys Arg Lys Asn Ile Glu Pro Thr His Ala Pro Phe Ile Glu Lys 130 140 Ala Asn Phe Asp Phe Asp Val TTP Arg Lys Lys Tyr Met Arg TTP Met 210 220 ğ Leu Ser Ala Arg Trp Gin Gin Val Trp Leu Leu Ala Leu Glu Arg Gin 180 185 Met Pro Trp Ala Lys Gln His Gln Gln Arg Leu Glu Thr Ala Leu Ser Glu 50 56 Lys Leu Asn Asp Ala Leu Asp Arg Leu Glu Glu Leu Lys Glu Phe 195 200 205 Ě (i) SEQUENCE CHARACTERISTICS: Ile Leu Ser Gln Ser Glu Ala Lys Asn Pro Arg Ile Asn Gln 165 170 175 SEQUENCE DESCRIPTION: SEQ ID NO: 429: (A) LENGTH: 370 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear 5 2 Αla

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SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 24 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 431:

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Gly Asn Thr Gly Arg Trp Leu Leu

Met Glu Pro His Leu Arg Cys Arg Val Thr Arg Val Arg Gly Ser Leu 1 10 15

45 8 35 ઝ 25 20 2 5 Gly Xaa 370 Ile Phe Asp Ary Asp Gly Asp Gly Tyr Ile Asp Tyr Tyr Glu Phe Val 275Ala Ser Lys Phe Pro Thr Thr Lys 260 Asp Gln Asp Gly Lys Ile Thr Arg Gln Glu Phe Ile Asp Gly Ile Leu 245 250 255 Asn His Lys Lys Ser Arg Val Met Asp Phe Phe Arg Arg Ile Asp Lys 225 230 230 Tie Leu Arg Asn Arg Asp Gly Ser Arg Trp Trp Arg Met Asp Gly Leu 355 Phe Leu Gly Asn Gln Phe Gly Asp Ser Gln Gln Leu Arg Leu Val Arg 340 345 Asp Lys Ile Glu Asp Glu Val Thr Arg Gln Val Ala Gln Cys Lys Cys 305 310 315 Ala Ala Leu His Pro Asn Lys Asp Ala Tyr Arg Pro Thr Thr Asp Ala 290 295 300 Ala Lys Arg Leu Tyr Leu Arg Tyr Val Thr Phe Val Tyr Leu Asn Leu Phe
20 25 30 Met Asn Val Lys Thr Phe Ser Xaa Asp His Met His Phe Leu Cys Cys
1 10 15 2 2 INFORMATION FOR SEQ ID NO: 430: INFORMATION FOR SEQ ID NO: 431: $\boldsymbol{\varepsilon}$ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 430: SEQUENCE CHARACTERISTICS: Phe Gin Val Glu Gin Ile Gly Glu Asn Lys Tyr Arg Phe 325 330 335 (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 30 amino acids Leu Glu Met Thr Ala Val Ala Asp 265 270

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	20	Val Trp Ser Pro Ser Thr Ser Arg Leu Thr Arg Tyr Thr Ile Trp His 145	
(2) INFORM	(2) INFORMATION FOR SEQ ID NO: 432:	S Leu Gin Pro Pro Leu Gin Thr Thr Cys Ile Ile Leu Ser Arg His Xas 170 170	
Z \$	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 53 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 432:		
let His 1	Met His Tyr Leu Val Leu Gly Gly Leu Gly Val Phe Leu Phe Ser 1 15	(2) INFORMATION FOR SEQ ID NO: 434:	
rys Phe V	Phe Ala Phe Phe I	15 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 77 amino acids (B) TYPS: amino acid	
yr Leu G	Tyr Leu Glu Gly Met Gly Gly Ser Gly Asn Ary Glu Val Gly Gly Gly 15 40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 434:	
he Cys 1	Phe Cys Leu Phe 50	net hell Aug ups in production and and and and and and and and and an	
		Cys Ser Leu Phe Trp Leu Leu Val Glu Trp Phe Gly Thr Asn Ile Asp 25 20 20 30 30 30 30 30 30 30 30 30 30 30 30 30	
2) INFOR	(2) INFORMATION FOR SEQ ID NO: 433:	Arg Glu Ser Tyr Asp Ala 11e Gly Gly Pro Ser Trp Met Thr Ala Ser 15	
	(1) SEQUENCE CHANACTERISTICS: (A) LENGTH: 176 amino acids (B) TYPE manno acid	30 Ser Phe Cys Leu Ser Asn Ser Asn Ile Trp Ser Leu Glu Ile Ser Ser 50	
×	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 433:	Gly Ser Thr Ser Val Val His Ser Gln Gln Ala Met Asp	
et val S 1	Met Val Ser Lys Ala Leu Leu Arg Leu Val Ser Ala Val Asn Arg 1	35 %	
rg Met Ly	Arg Met Lys Leu Leu Leu Gly Ile Ala Leu Leu Ala Tyr Val Ala Ser 20 25	(2) INFORM	
al Trp G	Val Trp Gly Asn Phe Val Asn Met Arg Ser Ile Gln Glu Asn Gly Glu 35 40	40 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 32 amino acids (D) TYPE: amino acid	
eu Lys I 50	Leu Lys Ile Glu Ser Lys Ile Glu Glu Met Val Glu Pro Leu Arg Glu 50 51	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 435:	
ys Ile A 65	Lys Ile Arg Asp Leu Glu Lys Ser Phe Thr Gln Lys Tyr Pro Pro Val 65 80	Met Arg Ser Cys Glu ile Gin Leu Cys Val Trp Leu Leu Val Ser Ser 1 10 15	
ys Phe L	Lys Phe Leu Ser Glu Lys Asp Arg Lys Arg Ile Leu Ile Thr Gly Gly 85 85	His Val Asp Met Val Leu Gly Gly Ser Pro Ser Thr Leu 1yr Met Met 50 25 20 30	
la Gly P	Ala Gly Phe Val Gly Ser His Leu Thr Asp Lys Leu Met Mep Gly 100 100		
iis Glu V	His Glu Val Thr Val Val Asp Asn Phe Thr Gly Arg Lys Arg Asn 115	(2) INFORMATION FOR SEQ ID NO: 436:	
al Glu F 130	Val Glu His Trp Ile Gly His Glu Asn Phe Glu Leu Ile Asn His Asp 130 135 140	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids	

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Leu Pro Cys Thr Pro Gln Met Val Arg Gly Val Thr Gln Val Leu Arg $50 \ \ \, 55$ Glu Leu Ser Thr Asp Ser Ser Ala Arg Leu Leu Tyr His Glu 20 25 30 Gin Gly Lys Ile Ala Phe Ser Leu Met Phe Val Leu Lys Asp Leu Ser $20 \ \ 30$ Met Val Val Asn Ser Leu Cys Phe Leu Ser Leu Leu Leu Val Ile Leu 1 5 10 Met Pro Leu Cys Phe Phe Ser Phe Leu Cys Cys Txp Val Leu Val Phe 1 10 15 Glu Phe Gly Asp Gln 65 Met Asp Lys Gln Lys His Leu Glu Val Arg Arg Ser Val Phe Lys Ile 1 10 Lys Leu Ile (2) INFORMATION FOR SEQ ID NO: 438: Pro Thr (2) INFORMATION FOR SEQ ID NO: 437: Œ. (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 436: Œ (1) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS: Tle Phe Ser His Ser Ile Leu Leu Leu Leu Pro His His Val 35 40 45 SEQUENCE DESCRIPTION: SEQ ID NO: 437: SEQUENCE DESCRIPTION: SEQ ID NO: 438: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 19 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 69 amino acids (D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 439:

(1) SEQUENCE CHARACTERISTICS:

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 439:

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 442:

(A) LENGTH: 64 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 442:

(1) SEQUENCE CHARACTERISTICS:

(D) TOPOLOGY: linear

(A) LENGTH: 43 amino acids (B) TYPE: amino acid

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Pro Pro

Pro Cys Leu Ser Ser Pro Phe Pro Phe Ile Ser Val Pro 20 25 30

Leu

Phe Glu Ala Val Pro Ile Ser Val Ser Asp Gln Pro Ser Pro Xaa 35 40 45

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Leu Thr Thr Leu Leu 50 8

Met Trp Gln Val Arg Gly Leu Pro Pro Val Pro Leu Leu Leu Thr Met
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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 441:

(A) LENGTH: 53 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 441:

(i) SEQUENCE CHARACTERISTICS:

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ટ્ટ 20 5 5 S 5 Leu Leu Asn Pro Ser Arg Leu Ile Leu Tyr Met Ile Ser Ala Gly Ala Asp Ser $20 \ 25 \ 30$ Met Leu Leu Phe Pro Ser Leu Leu Phe Ala Ala Thr Tyr Asn Val Ala 1 10 15 Arg Val Met Val Asn Leu Asn Ile Leu Phe Xaa 35 Met Lys Phe Ser Leu Val Leu Leu Ile Lys Ile Ile Ser Phe Glu Arg 1 19 (2) INFORMATION FOR SEQ ID NO: 440: (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 440: 11e Phe Leu 20 (A) LENGTH: 33 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear Phe Pro Leu Ser Phe Leu Pro Asn Ile Trp Arg 25

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Met Ile Thr Ser Val Leu Val Phe Leu Ile Phe Phe Phe Pro Tyr Leu 1 5

Ser Leu Val Thr Leu Leu Gln Ala Arg Asn Leu Trp Val 11e His Arg 25

S

Ala Ala Leu Cys Glu Ser Gly Leu Phe His Trp Arg Lys Gly Ile Glu 35 9 Asn Gln Leu Glu Pro Met Tyr Phe Leu Pro His Gly Thr Leu Phe Leu 50

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(2) INFORMATION FOR SEQ ID NO: 443:

2

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 34 amino acids

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 443: (D) TOPOLOGY: linear

25

Leu Tyr Ser Cys Glu Pro Tyr Leu Ile Ile Leu Asn Ile Tyr Ser 5 Met

Gin Lys Ala Phe Tyr Phe Tyr Phe Bhe Glu Gly Ser Phe Ser Val Cys $20\ 20$

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Thr Leu

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(2) INFORMATION FOR SEQ ID NO: 444:

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 89 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 444:

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Met Arg Gln Arg Gln Ala Ala Cys Gln Pro Pro Pro Ser Arg Asn Gly

Leu Ala Gln Glu Cys Pro Pro His Ile Pro Ser Ser Phe Phe Leu Val 25 တ္ထ

Leu Leu Phe Ile Pro Trp Leu Ala Ser Leu Leu Ser Ser Pro Leu 35 45 ξŞ

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Leu Gln Gln Cys Arg Gln His Gln Val Leu Gln Glu Lys Asn Thr $$70\ \ \, 70\ \ \, 75\ \ \, 80$ Leu Val Ser Ile Ser Trp Asp Leu Gly Leu Lys Leu 55 Ę Asn Leu Leu I 50 رة 5

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Lys Lys Phe Asn Lys Lys Lys Lys Lys 85

(2) INFORMATION FOR SEQ ID NO: 445:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 350 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 445:

Met Asp Phe Ile Thr Ser Thr Ala Ile Leu Pro Leu Leu Phe Gly Cys 1 5 10 15

2

Phe Gly Leu Phe Arg Leu Leu Gln Trp Val Arg Gly Lys 20 20 Leu Gly Val

Ala Tyr Leu Arg Asn Ala Val Val 11e Thr Gly Ala Thr Ser Gly 35

20

Leu Gly Lys Glu Cys Ala Lys Val Phe Tyr Ala Ala Gly Ala Lys Leu 50 60

22

Val Leu Cys Gly Arg Asn Gly Gly Ala Leu Glu Glu Leu Ile Arg Glu 65 70 75 80

Leu Thr Ala Ser His Ala Thr Lys Val Gln Thr His Lys Pro Tyr Leu

33

Val Thr Phe Asp Leu Thr Asp Ser Gly Ala Ile Val Ala Ala Ala Ala Ala 100 Glu Ile Leu Gln Cys Phe Gly Tyr Val Asp Ile Leu Val Asn Asn Ala 115 33

Gly Ile Ser Tyr Arg Gly Thr Ile Met Asp Thr Thr Val Asp Val Asp 130 \$

i Leu Thr Lys 160 Lys Arg Val Met Glu Thr Asn Tyr Phe Gly Pro Val Ala 145 155 145

Ala Leu Leu Pro Ser Met Ile Lys Arg Arg Gln Gly His Ile Val Ala 175 Ile Ser Ser Ile Gin Gly Lys Met Ser Ile Pro Phe Arg Ser Ala Tyr 180

45

Ala Ala Ser Lys His Ala Thr Gln Ala Phe Phe Asp Cys Leu Arg Ala 195 ဂ္ဂ

Glu Met Glu Gln Tyr Glu 11e Glu Val Thr Val 11e Ser Pro Gly Tyr 210 Ile His Thr Asn Leu Ser Val Asn Ala Ile Thr Ala Asp Gly Ser Arg 225

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Tyr Gly Val Met Asp Thr Thr Thr Ala Gln Gly Arg Ser Pro Val Glu 245 255

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Met Ala Ser Ala Glu Leu Asp Tyr Thr Ile Glu Ile Pro Asp Gln Pro 1 5 15 Pro Gly Leu Phe 290 Glu Thr Leu Met Glu Ile Cys Leu Thr Ser Gly Lys Asp 345 350 INFORMATION FOR SEQ ID NO: 446: INFORMATION FOR SEQ ID NO: 447: E (1) SEQUENCE CHARACTERISTICS: ž (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 447: SEQUENCE CHARACTERISTICS: Thr Phe Ile Leu Glu Thr Glu Val Tyr Leu Asp Leu Ala 20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 446: (A) LENGTH: 49 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 278 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Glu Ala Ala Leu Leu Gly Leu Leu Thr Leu Gln Gly Thr Val Ala Phe 325 330 335 Gly Asn Pro Arg Thr Pro Ser Thr Leu Thr Ser Gln Gly Gln Gly Arg 305 310 315 Ala Tie Leu Ala Asp Leu Leu Pro Ser Leu Ala Val Tyr Leu Arg Thr Leu 275 280 285 Val Ala Gln Asp Val Leu Ala Ala Val Gly Lys Lys Lys Lys Asp Val 260 265 Phe Ser Leu Met Pro Pro Gly Pro Glu Lys Ser 295

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Heet Val Phe Leu Pro Arg Gly Val Val Val Ser Gly Gly Ala Ala Cys

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Leu Trp Leu

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The Glu Ala Arg Ala His Ser Arg Met Gly Leu Gly Leu Trp Pro Pro 35 $$40^{\circ}$$

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SS Trp Ser Gln Lys Asn Ser Pro Ser Pro Gly Gly Lys Glu Ala Glu
20 25 30

Arg Gln Pro Val Val Ile Leu Leu Gly Trp Gly Gly Cys Lys Asp 35 † 40 45

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Lys Asn Leu Ala Lys Tyr Ser Ala Ile Tyr His Lys Arg Gly Cys Ile 50 60 Val Ile Arg Tyr Thr Ala Pro Trp His Met Val Phe Phe Ser Glu Ser 65 70 75 80

Leu Phe Asp Tyr Glu Ile Glu Lys Glu Pro Leu Leu Phe His Val Phe
100 105 110 Leu Gly Ile Pro Ser Leu Arg Val Leu 85 Ala Gln Lys Leu Leu Glu Leu 90 95

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5 Ser Asn Gly Gly Val Met Leu Tyr Arg Tyr Val Leu Glu Leu Leu Gln 115 120

쿭 Arg Arg Phe Cys Arg Leu Arg Val Val Cly Thr Ile Phe Asp Ser 130 140

20 Ile Leu Glu Arg Arg Ala Ala Met Leu Arg Leu Leu Leu Leu Val Ala 165 170 175 Ala Pro Gly Asp Ser Asn Leu Val Gly Ala Leu Arg Ala Leu Ala Ala 145 150 150

25 Ala Xaa Phe His Thr His Phe Tyr Asp Arg Leu Gln Asp Ala Gly Ser 195 200 205 Phe Ala Leu Val Val Leu Phe His Val Leu Leu Ala Pro Ile Thr 180 185

30 Arg Trp Pro Glu Leu Tyr Leu Tyr Ser Arg Ala Asp Glu Val Val Leu 210 \$210\$

Leu Ala Arg Ser Val Asp Phe Val Ser Ser Ala His Val Ser His Leu 245 255 Ala Arg Asp Ile Glu Arg Met Val Glu Ala Arg Leu Ala Arg Arg Val 225 230 230

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6 Arg Asp Tyr Pro Thr Tyr Tyr Thr Ser Leu Cys Val Asp Phe Met Arg 260 265

Asn Cys Val Arg Cys Xaa 275

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(2) INFORMATION FOR SEQ ID NO: 448:

 $\widehat{\Xi}$ SEQUENCE CHARACTERISTICS (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 199 amino acids

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Met Ser Phe Ile Phe Asp Trp Ile Tyr Ser Gly Phe Ser Ser Val Leu 1 5 10 Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 448:

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8 Cln Phe Leu Cly Leu Tyr Lys Lys Thr Cly Lys Leu Val Phe Leu Gly
20 25 30

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Leu Asp Asn Ala Gly Lys Thr Thr Leu Leu His Met Leu Lys Asp Asp
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5 Arg Leu Gly Gln His Val Pro Thr Leu His Pro Thr Ser Glu Glu Leu
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The le Ala Gly Met The Phe The The Phe Asp Leu Gly Gly His Val

10 Gln Ala Arg Arg Val Trp Lys Asn Tyr Leu Pro Ala Ile Asn Gly Ile 85 90 95

Val Phe Leu Val Asp Cys Ala Asp His Glu Arg Leu Leu Glu Ser Lys 15 100 105 110

20 11e Leu Ile Leu Gly Agn Lys Ile Asp Arg Pro Glu Ala Ile Ser Glu 130 135

Glu Arg Leu Arg Glu Met Phe Gly Leu Tyr Gly Gln Thr Thr Gly Lys 145 150 150 Met Cys Ser Val Leu Lys Arg Gin Gly Tyr Gly Glu Gly Phe Arg Trp 30 185 180

Gly Ser Ile Ser Leu Lys Glu Leu Asn Ala Arg Pro Leu Glu Val Phe 175

Met Ala Gin Tyr ile Asp Xee 195 (2) INFORMATION FOR SEQ ID NO: 449:

35

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 256 emino acide
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 449.

45 Met Thr Leu Ser Arg Phe Ala Tyr Asn Gly Lys Arg Cys Pro Ser Ser
1 5 10 10
15 Tyr Asn Ile Leu Asp Asn Ser Lys Ile Ile Ser Glu Glu Cys Arg Lys
20 25 30

50 Glu Leu Thr Ala Leu Leu His His Tyr Tyr Pro Ile Glu Ile Asp Pro 35 His Arg Thr Val Lys Glu Lys Leu Pro His Met Val Glu Trp Trp Thr 50 50 50 50 Leu Leu Cys Glu Glu Lys Ile Glu Lys Phe Glu Ile 65 70 80

60 Ala Gin Val Val Arg Giu Ser Asn Ala Met Leu Arg Giu Gly Tyr Lys

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Val Phe His Pro Asn Ile His Ile Val Ser Asn Tyr Met Asp Phe Asn 130

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Glu Asp Gly Phe Leu Gln Gly Phe Lys Gly Gln Leu Ile His Thr Tyr 145 145

Ash Lys Asn Ser Ser Val Cys Glu Asn Xea Gly Tyr Phe Gln Gln Leu 165

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Glu Gly Lys Thr Asn Val Ile Leu Leu Gly Asp Ser Ile Gly Asp Leu 180 190

8

Thr Met Ala Asp Gly Val Pro Gly Val Gln Asn Ile Leu Lys Ile Gly
195
200
200
Phe Leu Asn Asp Lys Val Glu Glu Arg Arg Arg Xaa Arg Tyr Met Asp Ser
210
210
215

25 210 215 220 Tyr Asp Ile Val Leu Glu Lys Asp Glu Thr Leu Asp Val Val Asn Gly 225 235 240 30 Leu Leu Gin His Ile Leu Cys Gin Gly Val Gin Leu Giu Met Gin Gly 255

Pro Xaa 35 (2) INFORMATION FOR SEQ ID NO: 450:

6

(1) SEQUENCE CHANGLISHISHIS:

(A) LENGTH: 87 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 450;

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Met Ser His Val Leu Leu Cys Pro Ser Leu Ser Cys Ser Asn Leu Leu

1 5 10 15

Pro Pro Ser His Ser Leu Gly Thr Met Gly Ser Leu Ser Pro His Leu
50 26 30 30

Pro Pro Ser His Ser Leu Gly Thr Met Gly Ser Leu Ser Pro His Leu 20 25 30 30 Cys Gly His Thr Met Cys Pro Val Asn Pro Glu Leu Pro Leu Ser Ser 35 40 40 45

Arg Leu Thr Thr Asp Gln Pro Gln Pro Asp Ala Cys Ser Pro Thr Leu 50 60

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Leu Thr Leu Pro Leu Pro Ser Ser Phe Leu Pro His Ser Lys Pro Thr 65 75 78

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Xaa His Pro Cys Ser Pro 85

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INFORMATION FOR SEQ ID NO: 451:

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 315 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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£ SEQUENCE DESCRIPTION: SEQ ID NO: 451:

Met Phe Ser Ile Asn Pro Leu Glu Asn Leu Lys Val Tyr Ile Ser Ser 1 15

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Arg Pro Pro Leu Val Val Phe Met Ile Ser Val Xaa Pro Met Ala Ile 20 25 30

Ala Phe Leu Thr Leu Gly Tyr Phe Phe Lys Ile Lys Glu Ile Lys Ser \$35\$

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Leu Asp Leu Cys Val Ser Glu Asn Glu Thr Leu Lys His Leu Thr Asn 65 70 75 80

Pro Glu Met Ala Glu Asp Trp Asn Thr Phe Leu 50 55

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Arg Phe Asn

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5 Lys Val Ala Leu Ala Glu Ala Xaa 310 315

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Ser

Tyr Phe Leu i 275

Phe Val Met Val Ile Thr Met Phe Cye Tyr Ala 280 285

Ile Lys Gly Arg Pro Ser Lys Leu Arg Gln Ser Asn Pro Glu Phe 290 295

Cys Pro Glu i 305

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(2) INFORMATION FOR SEQ ID NO: 452:

(1) SEQUENCE CHARACTERISTICS

(A) LENGTH: 52 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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£ SEQUENCE DESCRIPTION: SEQ ID NO: 452:

Met Pro Gly Leu Ser Leu Ala Leu Leu Pro Phe Gly Pro Gly Cys Thr 1 15

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Glu Ala Leu His Ala Gly Cys Phe Pro Ala Phe Ala Ser Ala Thr Arg $20 \\ 20 \\ 25$

Val Asn Gly Glu Ala Ala Leu Ser Pro Gly Leu Cys Asp Pro Ile Ser . 35 40 45

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Val Pro Tyr Val 50

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8 INFORMATION FOR SEQ ID NO: 453:

(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 383 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO:

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Gly Val Phe Pro Val Thr Val Gln Pro Pro His Cys Val Pro Asp Thr 195 200 205

Cys Glu Gln Val Val Phe Thr Ala Cys Met Thr Leu Thr Ala Ser Pro 180 185

Thr Leu Pro Thr Ala Trp Ser Ser Asp Asp Cys Ala Leu His Gly His 165 170 175

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Ser Arg Asn Val Thr His Leu Tyr Ser Thr Ile Leu Gly His Gln 130 140

Ile Gly Leu Ser Gly Arg Glu Ala His Glu Glu Ile Asn Ile Thr Phe 145 150 150

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Ser Val

Ser Ile Thr Leu Thr Leu Asp Pro Leu Lys Pro Phe Gly Gly 115 120 125

Ser Thr Gln

Ser Pro Gln Ala Leu Glu Asp Ser Gly Pro Val Asn Ile 100 105

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Asp Thr Thr

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Pro Glu Ser Thr Met Thr Ser Gly Gln Ala Arg Ala 85 90 95

Met Ala Val Gly Gln Ile Met Thr Phe Gly Ser Pro Val Ile Gly Cys
1 10 15

Gly Phe Ile Ser Gly Trp Asn Leu Val Ser Met Cys Val Glu Tyr Val 20 25 30

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Per Feu 2 **3** Lys Val Tyr Gln Lys Thr Pro Ala Leu Ala Val Lys Ala 40 45

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qι Asp Thr Glu Pro Lys Pro Leu Glu Gly Thr His Leu Met Gly Val Lys 65 70 75 80 Leu Lys Glu Glu Glu Thr Glu Leu Lys Gln Leu Asn Leu His Lys $50 \\ $ $$

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Ile Val Pro Asp Asp Asp Arg Ser Leu Ile Asn Leu His Leu Met His 260 265

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Gly Ala Ile Gly Lyb Val Tyr His Ala Leu Asn Pro Lys Leu Thr Val 245 250 250

Ala Asn Thr Lys Tyr Ala Gin Asp Tyr Asn Pro 235 230 230

Phe Trp Cys Tyr Lys 240

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Ser Asn Ala Thr Leu Trp Tyr Lys Ile Phe Thr Thr Ala Arg Asp 210 215

Thr Ser 175 Asp Ser Asn Ile His Glu Leu Glu His Glu Glu Glu Pro Thr Cys Ala 95 Phe Arg Asp Gly Trp Val Ser 110 Leu Ala Phe Leu 125 Thr Gln Cly Leu Ser Gly Phe His Pro Gln Tyr Phe Asp Gly Ser Ile 145 Thr Gly Tyr Ala Tyr 140 Lys Met Trp Pha Gly Ser Ala Gly Leu Ile Ser Gly Leu Ala Gln Leu 180 Ser Cys Leu Ile Leu Cys Val Ile Ser Val Phe Met Pro Gly Ser Pro 195 Gin Gly Glu Ser Ile Thr Pro Thr Lys Ile Pro Glu Ile Thr Thr Glu 125 Phe Ile Pro Glu Ser Val Pro Ile Ile Ser Val Ser Leu Leu Phe Ala Gly 260 Leu Trp Ser Phe Asp Leu Thr Val Thr 280 Met Val Ile Leu Ala Pro Agn Pro Glu Ala Phe Gly Leu Val Leu 335 Ser Phe Val Ala Met Gly His Ile Met Tyr Phe Arg Phe 340 Lys Leu Phe Ala Cys Gly Pro Asp Ala 360 Ile Tyr Met Ser Asn Gly Ser Asn Ser Ala Asn Ile Val Pro Glu Thr 255 Leu Leu Gln Glu Asn Val Ile Glu Ser Glu Arg Gly Ile Ile Asn 290 Phe Ile Ser Val Val Xaa Ser Tyr Asn Trp Asn Asn Gly Asn Cys Ser Phe Tyr Leu Ala 165 Leu Asp Leu Ser Val Ser Pro Phe Glu Asp Ile Arg Ser Arg 210 Val Gln Asn Ser Met Asn Tyr Leu Leu Asp Leu Leu His $$310\$ Tyr Tyr Asn Gln Pro Val Phe Leu Ala Gly Met Gly 115 Phe Asp Cys Ile Thr 135 Asn Thr Glu Val Arg Lys Glu Asn Gln Ala 370 Ser Gin Met Ala Glu Pro Phe Arg 100 Agn Val Ile Ala Ala Arg Ile Gly Leu Gly Ala Gln Asn Thr Leu Gly 355 Thr Val Ser Val Met . ጟ Ser G1y 305 g Ile. Lya 9 2 22 6 15 ಜ 35 တ္တ 55 5

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(A) LENGTH: 186 amino acida (i) SEQUENCE CHARACTERISTICS

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TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 454:

Met Arg Ser Ile Gly Asn Lys Asn Thr Ile Leu Leu Gly Leu Gly Phe 1 5

Gin ile Leu Gin Leu Ala Trp Tyr Gly Phe Gly Ser Glu Pro Trp Met 2

Met Trp Ala Ala Gly Ala Val Ala Ala Met Ser Ser Ile Thr Phe Pro 35 Ala Val Ser Ala Leu Val Ser Arg Thr Ala Asp Ala Asp Gln Gln Gly 50 60

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9 g Gly Pro Ala Leu Tyr Gly Phe Ile Phe Tyr Ile Phe His Val Glu Leu Val Val Gln Gly Met Ile Thr Gly Ile Arg Gly Leu Cys Asn Gly 65 ន

Lys Glu Leu Pro Ile Thr Gly Thr Asp Leu Gly Thr Asn Thr Ser Pro 100 23

Pro Pro Phe Leu 125 Gln His His Phe Glu Gln Asn Ser Ile Ile Pro Gly 115 $$\rm 120$ 8

Trp Arg Lys His 160 Ser Ser 155 Asn Leu Ser Leu Arg Ser 150 녍 Pro Glu His ? 145 35

Leu Val Ala Leu Phe Ile 140

Gly Ala Cys Ser Val Leu Leu Ala Leu 130

Phe

Ser His Ser His Pro His Asn Thr Gln Ala Pro Gly Glu Ala 170 175 Asn Val 185 뀵 Leu Leu Gln Asp 180 Pro GJn Lya 6

Cys Gly

INFORMATION FOR SEQ ID NO: 455: 3

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SEQUENCE DESCRIPTION: SEQ ID NO: 455: (A) LENGTH: 163 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear XI)

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Met Leu Gln Thr Ser Asn Tyr Ser Leu Val Leu Ser Leu Gln Phe Leu 1 15 15 Tyr Asp Leu Phe Val Asn Ser Phe Ser Glu Leu Leu Gln 20 30 Leu Ser 25

Lys Thr Pro Val 11e Gln Leu Val Leu Phe 11e 11e Gln Asp 11e Ala S

(2) INFORMATION FOR SEQ ID NO: 454:

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Glu Ser Ser Pro Thr Glu Thr Ser Glu Gln Ile Arg Glu Lys 35 40 45Gln Phe Thr Lys Pro Pro Ser Leu Pro Leu Glu Pro Glu Pro Ala Val 20 25 30 Met Arg Ile Gln Val Phe Ile Leu Leu Leu Gly Ala Gly Gly Thr Ser 1 10 15 His Phe Tyr Gln Asp 145 궃 Met Ser Tyr Leu Ala Phe Leu Tyr Met Thr Phe Asp Phe Cys Cys Leu 1 15 (2) INFORMATION FOR SEQ ID NO: 457: Arg Arg Xaa (2) INFORMATION FOR SEQ ID NO: 456: (i) SEQUENCE CHARACTERISTICS: χĽ Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 457: SEQUENCE DESCRIPTION: SEQ ID NO: 456: (A) LENGTH: 46 emino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 105 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Ser Leu Trp Leu Arg Lys Glu Phe Met Gln Val 150 155 160

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20 25 5 5 Thr Asp Gly Leu Gln Met Leu 115 The Ile Ile Leu Thr Ala Val Tyr Phe Ala Leu Ser Ile Ser Leu His $95 \hspace{1.5cm} 95$ Val Phe Gln Ala Cly Leu Val Asn Leu Leu Phe His Lys Phe Lys Gly 65 70 75 Val Leu Phe Asn Ile Ile Ile Ile Phe Leu Met Phe Phe Asn Thr Phe $50\,$ Leu Val Trp Val Met Asn Leu Arg Trp Lys Asn Ser Asn Ser Phe Ile Trp $100\,$ Tyr Cys Tyr Phe Tyr Lys Arg Thr Ala Val Arg Leu Gly Asp Pro 130 140 Phe Val Phe Gln Arg Leu Ala Ala Val 120 125

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Ser Lys Ser Ser Ala Glu Ala Asp Gly Val 50 55

Leu Gln Pro Arg Arg 60

Phe Ala Val Ser Leu Ala Ala Lys Xaa 100

Ser Leu Leu Ile Phe Ser Phe Gln Lys Thr Glu Ala Lys Leu Ile Val 95

His Pro Ala Ser Leu Leu Ile Val Phe Ala Thr Ser Ile Ser Glu Ser 65 70 75 80

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(1) SEQUENCE CHARACTERISTICS

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INFORMATION FOR SEQ ID NO: 458:

(A) LENGTH: 70 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

£ SEQUENCE DESCRIPTION: SEQ ID NO: 458:

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Leu Leu Pro Val Ile Leu Ala Val Phe Pro Asp Ser Ala Gln Ile Val
20 25 30 Met Leu Pro Pro Phe Ser Leu Val Tyr Thr His Phe Leu Val Ala Ser 1 5 10

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Pro Leu Leu Lys Pro Ile Pro Arg Pro Gln Pro Glu Val Ile Phe Pro 35 40 45

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Ser Ser Glu Leu Leu Glu Oln Leu Leu Ser Val Gln Phe Val Trp Oln $50 \ \ \, 50$

Ala His Thr Val Ala Xaa 65 70

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ຣ INFORMATION FOR SEQ ID NO: 459:

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E SEQUENCE CHARACTERISTICS:

(A) LENGTH: 155 amino acids

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 459: (B) TYPE: amino acid
(D) TOPOLOGY: linear

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Met Ala Leu Leu Ser Val Leu Arg Val Leu Leu Gly Gly Phe Phe 1 5

Ala Leu Val Gly Leu Ala Lys Leu Ser Glu Glu Ile Ser Ala Pro Val 20 25 30

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Ser Glu Arg Met Asn Ala Leu Phe Val Gln Phe Ala Glu Val Phe Pro 35 40 45

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Thr Asp Thr His Ile Cys Val Cys Val Cys Ile Tyr Leu Ser Ser Val
35 40 45

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Ala Cys Ala Arg Ser Ile Phe Ile Phe Asp Glu Met Asp Lys Met His 175 Ala Gly Leu Ile Asp Ala Ile Lys Pro Phe Leu Asp Tyr Asp Leu 180 Val Asp Gly Val Ser Tyr Gln Lys Ala Met Phe Ile Phe Leu Ser Asn 195 Ala Gly Ala Glu Arg Ile Thr Asp Val Ala Leu Asp Phe Trp Arg Ser 210 Gly Lys Gln Arg Glu Asp lle Lys Leu Lys Asp Ile Glu His Ala Leu 215 215 Ser Val Ser Val Phe Asn Asn Lys Asn Ser Gly Phe Trp His Ser Ser 250 245 Leu lle Asp Arg Asn Leu lle Asp Tyr Phe Val Pro Phe Leu Pro Leu 260 $$260\,$ Glu Tyr Ly9 His Leu Ly9 Met Cy8 Ile Arg Val Glu Met Gln Ser Arg $_{\rm 275}$ Gly Tyr Glu Ile Asp Glu Asp Ile Val Ser Arg Val Ala Glu Glu Met 290 290 Phe Phe Pro Lys Glu Glu Arg Val Phe Ser Asp Lys Gly Cys Lys 310 Met Ile Leu Thr Leu Leu Ser Val Val Ser Thr Met Ala Ser 1 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 462: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 461 Lys Leu Asp Tyr Tyr Tyr Asp Asp 325 (A) LEWSTH: 14 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 5 emino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS INFORMATION FOR SEQ ID NO: 461: (2) INFORMATION FOR SEQ ID NO: 462: Leu Lys Cys Ile 5 Thr Val Phe Thr 고 3 3 5 Met 3 S 2 12 25 2 23 8 33 4 45 S 8 Leu Leu Ala Gly Leu Leu Leu Val Met Gly Pro 70 78 80 Leu Lys Val Phe Gly Tyr Gln Pro Asp Pro Leu Asn Tyr Gln Ile Ala 50 Pro Met Leu Gln Glu Ile Ser Asn Leu Phe Leu Ile Leu Leu Met Met 85 Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu Ser Thr Cys 100 Gly Phe Leu Leu Leu Asn Val Gly 120 Gin Leu Leu Ala Gin Thr Lys Lys Val Val Arg Pro Thr Arg Lys Lys 130 Ser 160 Lys Leu Gly Arg Ala Val Leu Gly Leu Leu Leu Leu Ala Pro Ser 10 Leu Ala Leu Ala Gly 30 Pro Arg Leu Tyr Cys Leu Phe Ala Glu 40 Leu Asp Asp Asp Leu Phe Cly Cln His Leu Ala Lys Lys Ile Ile Leu 65 78 Asn Ala Val Phe Gly Phe lle Asn Asn Pro Lys Pro Lys Lys Pro Leu 95 Val His Leu Phe Val Ala Thr Leu His Phe Pro His Ala Ser Asn Ile 130 Sp Thr Leu Ser Leu His Gly Trp Thr Gly Thr Gly Lys Asn Phe Val Ser 110 ጟ Cys Cys Gly Gln Lys Arg Sex Leu Sex Arg Glu Ala Leu Gln Lys 50 -60 Lys Ile Ile Ala Glu Asn Ile Tyr Glu Gly Gly Leu Asn Ser Asp 125 Leu Tyr Lys Asp Gln Leu Gln Leu Trp Ile Arg Gly Asn Val 150 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 460: Val Val Gln Ala Val Glu Pro Ile Ser Leu Gly 20 25 Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys Xaa 145 145 150 (A) LENGTH: 332 amino acids TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 460 lle Pro Ala Ile Val Cys Leu 115 Val Leu Thr Gly Tyr Ile Tyr Leu Glu Val Gly Phe 1 65 Met 142

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Αla Ala Gly Tyr Gly Phe Ala Phe Arg Tyr Cys Pro Ser Gly Lys His Val 145 150 150 Ala Glu Asn Ile Phe Val Asp Pro Glu Asp Gln Ser Gln Val Thr Leu $130\,$ Ş Ser Gly Arg Ser Cys Thr Arg Pro His Cys Trp Pro Ser Leu Pro Ala 50 55 Ala Leu Thr Tyr Glu Glu Lys Pro Pro Tyr Ala Met Leu Arg Asn Asn 260 265 Gln Lys Phe Val Asp Lys Pro Gly Pro Phe Val Gly Pro Cys Gly His 225 230 240 ř Ę Ile 145 Ala Leu Glu Phe Leu His Glu Asn Glu Tyr Val His Gly Asn Val Thr 115 120 125 Ala Trp Gly Gly Ala Phe Ser Arg Pro Trp Met Ser Ala Gln Ser Met
90
95 85 13 Ser Lys Ser Ser His Ser Asn Trp Met Pro Arg Met Gly Ala Cys Ser 20 25 Met Lys Leu His Pro Pro Pro Pro Ser Pro Val Thr Gln Asp His Arg
1 15 Trp Ile Arg Pro . C Tyr Val Glu Gly Ser Arg Ser Pro His Glu Gly Asp Leu Glu Phe $165\,$ ŝ Ser Trp Thr Asn Cys Leu Pro Xaa Xaa Glu Asp Ile Met Lys Gln Lys 210 225 Ser Met Asp Leu His Lys Gly Cys Gly Pro Ser Arg Arg Xaa Asp 180 185 Val Ser Val INFORMATION FOR SEQ ID NO: 463: Ser Leu Gly Tyr Cys Met Leu Lys Trp Leu Tyr Gly Phe Leu 195 200 205 Ala Glu Arg Ser Val Leu 100 (i) SEQUENCE CHARACTERISTICS: Arg Thr Ser Ser Ser 35 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 285 amino acids 뫔 Ser Glu Thr Leu Gln Lys Tyr Leu Lys Val Val Met 245 250 255 The Arg The Asn The Gly See Trp Cys Tyr Pro 70 75 80 40 Pro Gln Val Ala Cys Arg Leu Leu Asp 105 110 Pro Ser Leu Cys Lys Ser Thr 45

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8 55 8 25 8 Met His Thr Trp Tyr Asn Asp Arg Arg Gln Asn Cys His Cys Leu Leu 1 5 10 Pro Leu Leu Val Lys Cys Arg Gly Arg Leu Lys Gly Val Asn Ile 35 40 45 ອ Phe Phe Leu INFORMATION FOR SEQ ID NO: 466: £ Ĕ SEQUENCE CHARACTERISTICS: Ile Tyr Leu Arg Lys Ile Tyr Gln Val Val Pro His Val
20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 466: SEQUENCE DESCRIPTION: SEQ ID NO: 465: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 96 amino acids (D) TOPOLOGY: linear

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Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENOTH: 47 amino acids

35 (2) INFORMATION FOR SEQ ID NO: 465:

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Trp Gly Lys Arg Ile Ile Ser Glu His Cys Ser Ala Gln Ser Ser Xaa 65 70 75 80 Ą Tyr Cys Leu Leu Val Ala Asn Gln Ser Ile Phe Phe Pro Cys Leu 35 40 45 Glu Ser Gln Val Pro Leu Ala Leu Ser Arg Val Phe Ser Thr Ser His $20\ 25\ 30$ Met Thr Ser Pro Pro Pro His Gln Gly Tro Glu Gln Arg Gly Cys Gly 1 $$ 15 Ala Val Glu Arg Leu Leu Gly Val Arg Cys Thr Cys Pro Leu Ser 50 55

ξ. SEQUENCE DESCRIPTION: SEQ ID NO: 464:

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Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 80 amino acids

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Leu Glu Ala Leu Leu Gln Asp Leu Arg Val Ser Pro 275 28 각

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(2) INFORMATION FOR SEQ ID NO: 464:

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Met Glu Leu Val Leu Val Phe Leu Cys Ser Leu Leu Ala Pro Met Val 1 5 Leu Ala Ser Ala Ala Glu Lys Glu Lys Glu Met Asp Pro Phe His Tyr 20 25 Asp Tyr Gln Thr Leu Arg Ile Gly Gly Leu Val Phe Ala Val Val Leu 35 Asn Leu Ile Thr Ala Asn Ala Thr Glu Pro Gln Lys Ala Glu Asn Xaa 95 Leu Leu Ile Leu Ser Arg Arg Cys Lys Cys Ser 55 Asn Gin Lys Pro Arg Ala Pro Gly Asp Glu Glu Ala Gln Val Glu 75 75 80 116 Ser Val Gly 1 50 Phe A 62 2 2

INFORMATION FOR SEQ ID NO: 467 5 25

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 467; SEQUENCE CHARACTERISTICS:
(A) LENGTH: 399 entino acids
(B) TYPE: entino acid
(D) TOPOLOGY: linear 30

Met Ala Ser Gly Ala Asp Ser Lys Gly Asp Asp Leu Ser Thr Ala Ile 1 5 10 15 Leu Lys Gln Lys Asn Arg Fro Asn Arg Leu Ile Val Asp Glu Ala Ile 20 35

Asn Glu Asp Asn Ser Val Val Ser Leu Ser Gln Pro Lys Met Asp Glu 35 Leu Gln Leu Phe Arg Gly Asp Thr Val Leu Leu Lys Gly Lys Lys Arg $50\ \ \, 50$ 6

Ser Asp Glu Cys Ile Val Leu Ser Asp Asp Thr Cys 70 75 Arg Glu Ala Val 65 45

Asn Asn Leu Arg Val Arg Leu 90 Pro Asp Val Lys Tyr Gly Lys 110 Ser Ile Gln Pro Cys 105 Lys Ile Arg Met Asn Arg Val Val Arg Asp Val

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Leu Glu Ala Tyr Arg 140 ile Thr Gly Glu Gly 125 Arg Ile His Val Leu Pro Ile Asp Asp Thr Val 115 돲 Asn Leu Phe Glu Val Tyr Leu Lys Pro Tyr 130 55

Pro Ile Arg Lys Gly Asp Ile Phe Leu Val Arg Gly Gly Met Arg Ala 8

160 155 150

145

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Val Glu Phe Lys Val Val Glu Thr Asp Pro Ser Pro Tyr Cys Ile Val 165 Ala Pro Asp Thr Val Ile His Cys Glu Gly Glu Pro Ile Lys Arg Glu 180 180

Amp Glu Glu Ser Leu Asn Glu Val Gly Tyr Amp Amp Ile Gly Gly 200 Cys Arg Lys Gln Leu Ala Gln Ile Lys Glu Met Val Glu Leu Fro Leu 210 2

Leu Phe Lys Ala Ile Gly Val Lys Pro Pro Arg Gly 230 230 235 Thr Gly Lys Thr Leu Ile Ala Arg 250 . 255 Ile Leu Leu Tyr Gly Pro Pro Gly 245 Arg His Pro Ala L 225 2

Ala Val Ala Asn Glu Thr Gly Ala Phe Phe Leu Ile Asn Gly Pro 260 Glu Ser Glu Ser Asn Leu Arg Lys 285 Glu Ile Met Ser Lys Leu Ala Gly 275

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Glu Leu Asp Ala Ile Ala Pro Lys Arg Glu Lys Thr His Gly Glu Val 305 33

Ala Phe Glu Glu Ala Glu Lys Asn Ala Pro Ala Ile Ile Phe Ile Asp 290

Glu Arg Arg Ile Val Ser Gln Leu Leu Thr Leu Met Asp Gly Leu Lys 335 Gin Arg Ala His Val Ile Val Met Ala Ala Thr Asn Arg Pro Asn Ser 340

35

Ile Asp Pro Ala Leu Arg Arg Phe Gly Arg Phe Asp Arg Glu Val Asp 365 lle Gly lle Pro Asp Ala Thr Gly Arg Leu Glu lle Leu Gln lle His 370 5

Leu Ala Asp Asp Val Asp Leu Glu Gln Xaa 390 Thr Lys Asn Met Lys 385 45

INFORMATION FOR SEQ ID NO: 468 6

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SEQUENCE DESCRIPTION: SEQ ID NO: 468: ž:

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Thr Leu Val Arg Ser Phe Trp Ser Asp Met Met Asp Ser Ala Gln Ser 145 150 150 Glu Gln Leu Met Ser Leu Met Pro Lys 130 135 His Arg Ala Cys Gln Leu Thr Tyr Pro Leu His Thr Tyr Pro Lys Glu 50 Ala Ser Ala Glu Ala Phe Asp Ser Val Leu Gly Asp 35Pro Pro Leu Leu Leu Thr Met Ala Leu Ala Gly Gly Ser Gly Thr $20 \ \ 30$ Met Ala Ala Pro Lys Gly Ser Leu Trp Val Arg Thr Gln Leu Gly Leu 1 10 15 Met Leu Glu Pro Gly Ala Leu Pro Asn Leu Kaa Kaa Kaa Ser Leu Ser Lys 200 205 Ile Val Ile Pha Xaa Ser Lys Pro Arg Asn Pro Arg Tyr Ala Pro His 180 185 Phe Ile Thr Ser Ser Trp Thr Phe Tyr Leu Gln Ala Asp Asp Gly Lys 165 170 175 Cys His Leu Gly Cys Gln Asn Gln Leu Pro Phe Ala Glu Leu Arg Gln 115 120 125 Glu Ser Gln Phe Val Asp Asp Gly Ile Asp Leu Asn Arg Thr Lys Leu Glu Cys 85 90 Glu Glu Leu Tyr Ala Cys Gln Arg Gly Cys Arg Leu Phe Ser Ile Cys 65 70 75 Ser Gly Trp Ile Leu Thr Thr Leu Val Leu Ser Val Met Val Leu
255
255 Leu Glu Asp, Gly Glu Ser Asp Gly Phe Leu Arg Cys Leu Ser Leu Asn 225 230 230 Ser Xaa Xaa Ser Xaa Met Arg Asn Ser Gln Ala His Arg Asn Phe 210 220 Ê Ala Ě SEQUENCE CHARACTERISTICS: Cys Thr Glu Ala Tyr Ser Gln Ser Asp Glu Gln Tyr Ala 100 105 110 SEQUENCE DESCRIPTION: SEQ ID NO: 469: (A) LENGTH: 273 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Met His Leu Leu Phe Pro Leu 140 o Thr Ala Ser (

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Leu Trp Ile Cys Cys Ala Thr Cys Cys Tyr Thr Leu Leu Asp Ala Val 260

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SEQUENCE CHARACTERISTICS

acids

(A) LENGTH: 192 amino (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 470:

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(2) INFORMATION FOR SEQ ID NO: 470:

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(2)

INFORMATION FOR SEQ ID NO: 469:

ઝ 20 30 82 3 8 8 Val Leu Thr J 145 Met Met Val Leu Ser Leu Gly Ile Ile Leu Ala Ser Ala Ser Phe Ser 1 10 15 Val Lys Gln Ala Thr Leu Asn Pro Ala 85 The Glu Lys Arg Leu The Lys Leu Leu Val His Ser Ser Leu Val Gly S0 $$ Pro Asn Phe Thr Gln Val Thr Ser Thr Leu Leu 25 Gly Ser Val Leu Phe Leu Pro His Ser Tyr Ile Gly Asn Ser Gly Met 165 170 175 Ser Ile Leu Ser Ala Leu Ser Ala Leu Val Gly Phe Ile Ile Leu Ser 65 70 75 80 Phe Ile Gly Pro Phe Phe Phe Ile Ile Ser Gly Ser Leu Ser Ile Ala 35 $40\,$ Xaa Leu Ser Leu Met Leu Ile Cys Thr Leu Leu Glu Phe Cys Leu Ala 130 140 Ser Leu Tyr Thr Thr Asp Cys Tyr Thr Ala Lys Ala Ser Leu Ala Gly
115 120 125 Lys Asn Asn Ile Pro Thr Arg Ser Tyr Val Ser Tyr Phe Tyr His Asp 100 105 Ser ž Ala Val Leu I 150 Met Thr His Asp Cys Gly Tyr Glu Glu Leu Leu Thr Ser 180 190 Arg Trp Lys Gln Ala Tyr Ser Asp Phe Pro 155 Ser Leu Gln Cys Glu Leu Asp 90 95 Asn Ser Ala Tyr Pro 30

636

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INFORMATION FOR SEQ ID NO: 471:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 234 amino acids
(B) TYPE: amino acid

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Met Arg Lys Thr Arg Leu Trp Gly Leu Leu Trp Met Leu Phe Val Ser 1 5 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 471:

(D) TOPOLOGY: linear

Glu Leu Arg Ala Ala Thr Lys Leu Thr Glu Glu Lys Tyr Glu Leu Lys 20

Glu Gly Gln Thr Leu Asp Val Lys Cys Asp Tyr Thr Leu Glu Lys Phe 35 2

Ala Ser Ser Gin Lys Ala Trp Gin Ile Ile Arg Asp Gly Glu Met Pro S0 60

Lys Thr Leu Ala Cys Thr Glu Arg Pro Sar Lys Asn Ser His Pro Val 65

2

Gln Val Gly Arg Ile Ile Leu Glu Asp Tyr His Asp His Gly Leu Leu 95

2

Arg Val Arg Met Val Asn Leu Gln Val Glu Asp Ser Gly Leu Tyr Gln 100

Cys Val Ile Tyr Gln Pro Pro Lys Glu Pro His Met Leu Phe Asp Arg 115 23

Ile Arg Leu Val Val Thr Lys Gly Phe Ser Gly Thr Pro Gly Ser Asn 130

Glu Asn Ser Thr Gln Asn Val Tyr Lys Ile Pro Pro Thr Thr Lys 145 8

Ala Leu Cys Pro Leu Tyr Thr Ser Pro Arg Thr Val Thr Gln Ala Pro 175 Pro Lys Ser Thr Ala Asp Val Ser Thr Pro Asp Ser Glu Ile Asn Leu 180 35

Thr Asn Val Thr Asp ile Ile Arg Val Pro Val Phe Asn Ile Val Ile 200 8

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Phe Ala Val Thr Leu Arg Ser Phe Val Pro 225

(2) INFORMATION FOR SEQ ID NO: 472:

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SEQUENCE CHARACTERISTICS:
(A) LENGTH: 105 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 472.

Met Leu His Ile Leu Pro Leu Lys Ser Tyr Asp Phe Pro His Phe Ser 1 $$10\,$

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Leu Phe Phe Phe Cys Ser Val Leu Trp Thr Phe Ser Asp Met His S

Arg Ser Gly Glu Asp Gly Pro Trp Thr Pro Cys Val His His Leu Ala 50 60

Phe Ser Pro Val Leu Phe Ile Glu Asn Pro Arg His Tyr Ala Asn Ala 95 Ala Ser Leu Ile Ser Tyr Gly Gln Pro Gly Phe Ile Cys Ile Ser Leu 65 70 75 2 2

Thr Val Thr Thr Leu Gly Asp Trp Xea 100

(2) INFORMATION FOR SEQ ID NO: 473:

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(A) LEWITH: 32 amino acids (B) TYPE: amino acid (i) SEQUENCE CHARACTERISTICS:

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 473: (D) TOPOLOGY: linear

Met Val Phe Leu Lys Tyr Arg Phe Leu Phe Phe Leu Val Phe Leu Ala $1 \ \ \, 1$ Asn Cys lle Tyr Ser Leu His Tyr Lys Pro Ser Leu Met Tyr Pro Lys 20 2

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(2) INFORMATION FOR SEQ ID NO: 474: \$

(A) LENGTH: 571 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 474:

Met Ala Leu Ser Arg Gly Leu Pro Arg Glu Leu Ala Glu Ala 1 1 5 Gly Gly Arg Val Leu Val Val Gly Ala Gly Gly Ile Gly Cya Glu Leu 20 ಜ

Leu Lys Asn Leu Val Leu Thx Gly Phe Ser His Ile Asp Leu Ile Asp . 35 25

Leu Asp Thr Ile Asp Val Ser Asn Leu Asn Arg Gln Phe Leu Phe Gln 50 55 8

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Leu Gly Leu 305 Asn Met Lys Ser Arg Phe Asp Ile Lys Ser Met Ala Gly Asn Ile Ile 370 , 375 , 380 Asp Phe Val Thr Ser Ala Ala Asn Leu Arg Met His Ile Phe Ser Met 355 360 365 Gly Asp Gly Ala Glu Leu Ile Trp Asp Lys Asp 340 345 Ē Ser Gin Gly Glu Glu Thr Asn Ala Ser Asp Gin Gin Asn Glu Pro Gin 290 100 ij Lys Glu Trp Ala Lys Ser Thr Gly Tyr Asp Pro Val Lys Leu Phe Thr $245\ 250\ 255$ Arg Ala Arg Ala Ser Asn Glu Asp Gly Asp Ile Lys Arg Ile Ser Thr 225 230 235 Asp Arg Ala Asp Pro Glu Ala Ala Trp Glu Pro Thr Glu Ala Glu Ala 210 215 220 Ŀγ Phe Asn Gln Leu Phe Gly Glu Glu Asp Ala Asp Gln Glu Val Ser Pro 195 200 205 Leu Gly Gln Val Thr Thr Ile Lys Lys Gly Val Thr Glu Cys Tyr Glu 145 150 150 155 Asn Thr Pro CY8 His Cys Leu Ala Ala Asp Val 130 Met Asn Ala Leu Asp Asn Arg Ala Ala Arg Asn His Val Asn Arg Met 115 120 125 Met Asn Pro Asp Tyr Asn Val Glu Phe Phe Arg Gln Phe Ile Leu Val 105 Leu Gln Phe Tyr Pro Lys Ala Asn Ile Val Ala Tyr His Asp 85 Lys Lys His Val Gly Arg Ser Lys Ala Gln Val Ala Lys Glu Ser Val 65 70 75 80 Phe ₽¥ Leu Phe Lys Asp Asp Ile Arg Tyr Leu Leu Thr Met Asp Lys Leu 260 265 Ser Lys Ser Ile Glu Thr Leu 325 Lys Arg Lys Pro Pro Val Pro Leu Asp Trp Ala Glu Val Gln 275 280 285 Pro Lys Pro Thr Gln Arg Thr Phe Pro Gly Cys Thr Ile Arg 165 170 Lys Asp Gln Gln Val Leu Asp Val Lys Ser Tyr Ala Arg 310 315 Ser Glu Pro Ile His Cys Ile Val Trp Ala Lys Tyr Leu 180 185 190 Pro Leu Ile Glu Ser Gly Thr Ala Gly Tyr 135 Arg Val His Leu Ala Glu Lys 330 335 Ąsp Pro Ser Ala Met 350 95

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2 INFORMATION FOR SEQ ID NO: 475:

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Asp Ala Ala Lys Ser Ile Thr Asn Gly Gln Xaa 565 570

Glu Val Val Gly Asp Ala Pro Glu Lys Val Gly Xaa Lys Gln Ala 545 550 555

560 560

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Leu Ile Asn Ile Leu 530

His

Ser Glu Asp Leu Gly Lys Asp Val Glu Phe 535 540

23

Asn Gly Ser Arg Leu Gln Ala Asp Asp Phe Leu Gln Asp Tyr Thr Leu 515 525

Glu Thr Glu Ala Asn Asn His Lys Lys Leu Ser Glu Phe Gly Ile Arg 500 505

Gln Ile Glu Asp Gly Lys Gly Thr Ile Leu Ile Ser Ser Glu Glu Gly 495

Gln Asp Lys Ile Val Lys Glu Lys Phe Ala Met Val Ala Pro Asp Val 465 470 475

Glu Val Thr Val Arg Leu Asn Val His Lys Val Thr Val Leu Thr Leu 450

Ala Leu Asp Pro Pro Asn Pro Asn Cys Tyr Val Cys Ala Ser Lys 435 440 445

Pro

Phe Leu Asn Lys Gln Pro Asn Pro Arg Lys Lys Leu Leu Val Pro Cys 420 425

Glu Gly Leu Lys Ile Leu Ser Gly Lys Ile Asp Gln Cys Arg Thr Ile 415

Pro Ala Ile Ala Thr Thr Asn Ala Val Ile Ala Gly Leu Ile Val Leu 385 390 390

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(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 312 amino acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

ž SEQUENCE DESCRIPTION: SEQ ID NO: 475:

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Met Gln Val Val Thr Cys Leu Thr Arg Asp Ser Tyr Leu Thr His Cys
1 10 15

Phe Leu Gln His Leu Met Val Val Leu Ser Ser Leu Glu Arg Thr Pro 20 25 30

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Thr Thr Gly Lys Met Glu Asn Tyr Glu Leu Ile His Ser Ser Arg Val 50 60 Ser Pro Glu Pro Val Asp Lys Asp 35 Phe Tyr Ser Glu Phe Gly Asn Lys 45

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Lys Phe Thr Tyr Pro Ser Glu Glu Glu Ile Gly Asp Leu Thr Phe Thr 65 70 75

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Val Ala Gln Lys Met Ala Glu Pro Glu Lys Ala Pro Ala Leu Ser Ile 95

Leu Leu Tyr Val Gln Ala Phe Gln Val Gly Het Pro Pro Pro Gly Cys \$100\$

Pro Leu Arg Pro Lys Thr Leu Lau Leu Thr Ser Ser Glu 126 Cys Arg Gly 115

9

Ile Phe Leu Leu Asp Glu Asp Cys Val His Tyr Pro Leu Pro Glu Phe 130

Ala Lys Glu Pro Pro Gln Arg Asp Arg Tyr Arg Leu Asp Asp Gly Arg 145 $$150\,$ 15

Leu Met Gly Tyr Gln Thr Tyr Pro 170 Arg Val Arg Asp Leu Asp Arg Val 165

Gin Pro Ser Pro Ser Ser Ser Het Thr Cys Lys Val Het Thr Ser Trp 180 2

Ala Val Ser Pro Trp Thr Thr Leu Gly Arg Cys Gln Val Ala Arg Leu 205

Glu Pro Ala Arg Ala Val Lys Ser Ser Gly Arg Cys Leu Ser Pro Val 210 25

Leu Arg Ala Glu Arg Ser Ser Arg Cys Trp Leu Ala Ser Gly Arg 225 235 Pro Cys Val Ala Val Ser Cys Leu Ser Ser Pro Ala Ser Pro Gly 250 255 2

His Ser Gln Pro Val Val Ser Ser Leu Thr Pro Thr Gly Ala Gly Gln 260 35

Gin Ala Phe Val Phe Ser Lys Asn Val Leu Ser Ser Leu Trp Tyr Leu 275 Leu Thr Val Leu Ala Glu Asn Val Asn Met Cys Val Cys Cys Val 290 Asn

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Asn Ser Phe Ser Cys Trp Glu Xaa 305 45

INFORMATION FOR SEQ ID NO: 476: 3 တ္တ

(A) LENGTH: 329 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Ala Gin His His Leu Trp Ile Leu Leu Leu Cys Leu Gin Thr Trp \$1\$(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 476;

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Pro Glu Ala Ala Gly Lys Asp Ser Glu Ile Phe Thr Val Asn Gly Ile 8

Leu Gly Glu Ser Val Thr Phe Pro Val Agn Ile Glu Glu Pro Arg Gln 35 Ser Asp Leu Arg Met Glu Asp Ala Gly Asp Tyr Lys Ala Asp Ile Asn 100 Val Lys lle lle Ala Trp Thr Ser Lys Thr Ser Val Ala Tyr Val Thr 50 60 Pro Gly Asp Ser Glu Thr Ala Pro Val Val Thr Val Thr His Arg Asn 65 78 Tyr Tyr Glu Arg Ile His Ala Leu Gly Pro Asn Tyr Asn Leu Val Ile 85 The Gin Ala Asp Pro Tyr The The The Lys Arg Tyr Asn Leu Gin Ile 115 2 15 8

Tyr Arg Arg Leu Gly Lys Pro Lys Ile Thr Gln Ser Leu Met Ala Ser 130

Val Asn Ser Thr Cys Asn Val Thr Leu Thr Cys Ser Val Glu Lys Glu 145 23

Glu Lys Asn Val Thr Tyr Asn Trp Ser Pro Leu Gly Glu Glu Gly Asn Val Leu Gln Ile Phe Gln Thr Pro Glu Asp Gln Glu Leu Thr Tyr Thr Cys Thr Ala Gin Asn Pro Val Ser Asn Asn Ser Asp Ser Ile Ser Ala ಜ

Arg Gln Leu Cys Ala Asp Ile Ala Met Gly Phe Arg Thr His His Thr 210 35

Gly Leu Leu Ser Val Leu Ala Met Phe Phe Leu Leu Val Leu Ile Leu 235 Ser Ser Val Phe Leu Phe Arg Leu Phe Lys Arg Arg Gln Asp Ala Ala 255 수

Ser Lys Lys Thr Ile Tyr Thr Tyr Ile Met Ala Ser Arg Aan Thr Gln 260 ž Pro Ala Glu Ser Arg Ile Tyr Asp Glu Ile Leu Gln Ser Lys Val 280 45

Pro Ser Lys Glu Glu Pro Val Asn Thr Val Tyr Ser Glu Val Gln Phe 290 Ala Asp Lys Met Gly Lys Ala Ser Thr Gln Asp Ser Lys Pro Pro Gly 305 ೱ

Ser Tyr Glu Ile Val Ile Thr Ser

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55 8 Tyr Met Gln Lys Thr Leu Arg Gln Cys Gln Glu Gln Arg Gln Cys His 115 120 125 Leu Leu Ala Phe Tyr Val Asp Arg Val Phe Lys Asp His Gln Glu Pro 85 90 95 Gln Ala Lys Asp Thr Phe Pro Asn Val Thr Ile Leu Ser Thr Leu Glu 50 55 Asp Met His His Ile Glu Glu Ser Phe Gln Glu Ile Lys Arg Ala Ile \$35\$Leu Cys Ser Val Asp Asn His Gly Leu Arg Arg Cys Leu Ile Ser Thr $20 \ 30$ Asp Val Phe Leu Ala Trp Ile Asn Lys Asn His Glu Val Met Ser Ser 165 170 Cys Arg Gln Glu Ala Th
x Asn Ala Th
x Arg Val Ile His Asp Asn Tyr 130 140 Asn Pro Lys Ile Leu Arg Lys Ile Ser Ser Ile Ala Asn Ser Phe Leu 100 105 Thr Leu Gin Ile Ile Lys Pro Leu Asp Val Cys Cys Val Thr Lys Asn 65 70 75 80 Asp Gln Leu Glu Val His Ala Ala Ala Ile Lys Ser Leu Gly Glu Leu 145 150 150 Asp Thr Ala Ile Arg Val Ala Leu Ala Val Ala Val Leu Lys Thr Val 1 5 10 2 Ala Xaa Ile Leu Gly Leu Leu Cys Leu Leu Cys Gly Gly Gly Gly Gly Lys
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25 INFORMATION FOR SEQ ID NO: 478: (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 52 amino acids (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 478: (B) TYPE: amino acid

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Met Lys Leu Gin Cys Val Ser Leu Trp Leu Leu Gly Thr Ile Leu Ile 1 15 Ξ Œ. SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 477: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 178 amino acids

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(2) INFORMATION FOR SEQ ID NO: 477:

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Val Ala Gly Arg Gln Ala Val Thr Ser Asp Gln Gln Ser Val Gly Arg
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Arg Asp Val Tyr 50

5 2 INFORMATION FOR SEQ ID NO: 479:

(i) SEQUENCE CHARACTERISTICS

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 479: (A) LENGTH: 62 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

15

20 Met Gln Lys Lys Asn Ser Leu Phe Phe Phe Phe Ala Phe Tyr Tyr Glu 10

Asn Lys Thr Asn Ala Pro Gly Glu Gly Ser Met Ile Thr Arg Asn Ile 20 25

23 Ala Ile Asn Lys Leu Asn Tyr Leu His Trp Thr His Phe Gln $_{50}$ Lys Glu Tyr Phe Leu Pro 35 Phe Leu Phe Cys Cys Val Glu Ala Ser Ile 40 45

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(2) INFORMATION FOR SEQ ID NO: 480:

(i) SEQUENCE CHARACTERISTICS: <u>£</u> SEQUENCE DESCRIPTION: SEQ ID NO: 480: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 27 amino acids

35

6 Met Pro Gly Leu Ser Leu Ile Leu Thr Val Thr Leu Leu Ala Val Ser 1 15

Asp Ser Ala Ala Thr Cys Ile Val Ala Lys Gly
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2 INFORMATION FOR SEQ ID NO: 481:

8 (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 339 amino acids (D) TOPOLOGY: linear

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SS Met Ser Gly Pro Asp Val Glu Thr Pro Ser Ala Ile Gln Ile Cys Arg 1 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 481:

The Met Arg Pro Asp Asp Ala Asn Val Ala Gly Asn Val His Gly Gly $20 \ 20$

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Thr Ile Leu Lys Met Ile Glu Glu Ala Gly Ala Ile Ile Ser Thr Arg 35

His Cys Asn Ser Gln Asn Gly Glu Arg Cys Val Ala Ala Leu Ala Arg S

Phe Leu Ser Pro Met Cys Ile Gly Glu Val Ala 70 75 80 Val Glu Arg Thr Asp 65

9

His Val Ser Ala Glu Ile Thr Tyr Thr Ser Lys His Ser Val Glu Val 95

Gin Val Ann Val Met Ser Glu Asn Ile Leu Thr Gly Ala Lys Lys Leu 100

2

Thr Asn Lys Ala Thr Leu Trp Tyr Val Pro Leu Ser Leu Lys Asn Val 115

Lys Val Leu Glu Val Pro Pro Val Val Tyr Ser Arg Xea Glu Gln 130 Agp 2

Glu Glu Glu Gly Arg Lys Arg Tyr Glu Ala Gln Lys Leu Glu Arg Met 160 Glu Thr Lys Trp Arg Asn Gly Asp Ile Val Gln Pro Val Leu Asn Pro 175

22

Glu Pro Asn Thr Val Ser Tyr Ser Gln Ser Ser Leu Ile His Leu Val 180

8

Gly Pro Ser Asp Cys Thr Leu His Gly Phe Val His Gly Gly Val Thr 195

Lys Leu Met Asp Glu Val Ala Gly Ile Val Ala Ala Arg His Cys 210 Lys Thr Asn Ile Val Thr Ala Ser Val Asp Ala Ile Asn Phe His Asp 225 Met 35

Lys Ile Arg Lys Gly Cys Val Ile Thr Ile Ser Gly Arg Met Thr Phe 255 6

Thr Ser Asn Lys Ser Met Glu Ile Glu Val Leu Val Asp Ala Asp Pro 260 45 Val Val Asp Ser Ser Gln Lys Arg Tyr Arg Ala Ala Ser Ala Phe Phe 275

Thr Tyr Val Ser Leu Ser Gin Glu Gly Arg Ser Leu Pro Val Pro Gln 290 S

Leu Val Pro Glu Thr Glu Asp Glu Lys Lys Arg Phe Glu Glu Glu Gly Lys 320 Gly Arg Tyr Leu Gln Met Lys Ala Lys Xaa Gln Gly His Ala Xaa Xaa 330 55

Gln Pro Xaa පි

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(2) INFORMATION FOR SEQ ID NO: 482:

(A) LENGTH: 32 amino acids (i) SEQUENCE CHARACTERISTICS:

TYPE: amino acid

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Met Leu Asn Ser Asn Ile Asn Asp Leu Leu Met Val Thr Tyr Leu Ala $_{\rm 1}$ $_{\rm 1}$ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 482:

Asn Leu Thr Gin Ser Gin Ile Ala Leu Asn Giu Lys Leu Val Asn Leu 20 15

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(2) INFORMATION FOR SEQ ID NO: 483:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 48 amino acids

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 483:

3

Met Arg Glu Thr Ser Ile Arg Val Leu Leu Met Leu Pro Ala Leu Glu 1 10 11 15

Ser Thr Ser Gly Leu Ser Ala Phe Met Gly Leu Gly Thr Arg Ile Gly 25

35

Cys Phe Lys Thr Ile Thr Cys Trp Pro Thr Ser Leu Thr Gln Arg Xaa 35

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(2) INFORMATION FOR SEQ ID NO: 484: 45 SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 484:

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Met Tyr Met Tyr Ser Leu Asn Val Phe Leu Ser Phe Ile Phe Leu Ala 1 $_{\rm 1}$ 25

Leu Val Phe Lys Cys Val His Val Cys Gln Gly Ala Asn Ala Phe Leu 20

Phe Leu Lys Leu Val Phe 35

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SEQUENCE CHARACTERISTICS

INFORMATION FOR SEQ ID NO: 485:

30 20 5 5 .4 40 ၓ 25 S 50 Leu Val Phe Ala Trp Glu Phe Phe Ser Glu Asp Thr Pro 50 55 60 Gln Glu Cys Thr Glu Lys Phe Ala Lys Leu Leu Val Gln Leu Ile Ser \$45\$Val Cys Glu Met Phe Leu Phe Phe Leu Met Thr Gln Lys Leu Ile Trp $20 \ 30$ Met Gly Leu Arg Leu Ile Cys Leu Glu Leu Thr Met Val Lys Ala Leu 1 5 Thr Lys Asn Gln Trp Leu Leu Thr Pro Ser Arg Glu Tyr Ala Thr Lys $35 \ \ 40 \ \ \ 45$ Met Leu Ala Ala Arg Leu Val Cys Leu Arg Thr Leu Pro Ser Arg Val The Arg Ile Gly Ile Arg Arg Gly Arg The Gly Gln Glu Leu Lys Glu $50 \ \mbox{50}$ Ala val Ile Trp Pro Gln Tyr Val Lys Asp Arg Ile His Ser Thr Tyr 115 120 125 Gly Arg Trp Phe Val Ala Gly Gly Ala Ala Val Gly Leu Gly Ala Leu 85 90 95 Ala Ala Leu Glu Pro Ser Met Glu Lys Ile 65 70 Phe His Pro Ala Phe Thr Lys Ala Ser Pro Val Val Lys Asn Ser Ile 20 30 (2) INFORMATION FOR SEQ ID NO: 486: Met Tyr Leu Ala Gly Ser Ile Gly Leu Thr Ala Leu Ser Ala Ile Ala 130 $_{\dagger}$ 135 ş Tyr Tyr Gly Leu Gly Leu Ξ £ (X SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 486: SEQUENCE DESCRIPTION: SEQ ID NO: 485: (B) TYPE: amino acid (A) LENGTH: 346 amino acids Ser Asn Glu Ile Gly Ala Ile Glu Lys 105 110 Phe Lys Ile Asp Gln Met 75 80

> 20 5 ဗ 25 5 $\frac{3}{5}$ Trp Leu Leu His Ser Gly Val Met Gly Ala Val Val Ala Pro Leu Thr 195 200 205 Ile Val Gly Gly Leu Ser Thr Val Ala Met Cys Ala Pro Ser Glu Lys 225 230 230 235 Ile Leu Gly Gly Pro Leu Leu Ile Arg Ala Ala Trp Tyr Thr Ala Gly 210 215 220 Val Arg Ser Ile Pro Tyr Asp Gln Ser Pro Gly Pro Lys His Leu Ala 180 185 Val Thr Ile Gly Val Thr Phe Ala Ala Met Val Gly Ala Gly Mat Leu 165 170 Ile Ser Arg Thr Pro Val Leu Met Asn Phe Met Met Arg Gly Ser Trp 145 150 Ser Pro Met Tyr Gly Val Gln Lys Tyr Asp Pro Ile Asn Ser Met Leu 305 310 310 Ala Thr Leu Tyr Ser Val Ala Met Tyr Gly Gly Leu Val Leu Phe Ser 275 280 285 Leu Ala Thr Gly Gly Asn Arg Lys Lys Xaa 340 345 Ser Ile Tyr Met Asp Thr Leu Asn Ile Phe Met Arg Val Ala Thr Met 325 330 335 Met. Val Phe Leu Asn Met Gly Ala Pro Leu Gly Val Gly Leu Gly Leu Val Phe 245 250 255 Phe Leu 290 Ser Ser Leu Gly Ser Met 260 Per. Tyr Asp Thr Gln Lys Val Ile Lys Arg Ala Glu Val $295\,$ Phe Leu Pro Pro Thr Thr Val Ala Gly 265 270

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SEQUENCE DESCRIPTION: SEQ ID NO: 487:

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Met Glu Glu Val Leu Leu Gly Leu Lys Asp Arg Glu Gly Tyr Thr 1 5 10

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(2) INFORMATION FOR SEQ ID NO: 487:

(1) SEQUENCE CHARACTERISTICS

(A) LEWOTH: 237 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

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Phe Trp Asn Asp Cys Ile Ser Ser Gly Leu Arg Gly Cys Met Leu 20 25 30

Ile Glu Leu Ala Leu Arg Gly Arg Leu Gln Leu Glu Ala Cys Gly Met
35 40 45

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Arg Arg Lys Ser Leu Leu Thr Arg Lys Val Ile Cys Lys Ser Asp Ala 50 55 60

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Ser Gly 95 Pro Thr Gly Asp Val Leu Leu Asp Glu Ala Leu Lys His Val Lys Glu 65 75 78 80 Glu Thr Trp Asn Pro Leu Lys Leu His Tyr Gln Leu Arg Asn Val Arg 100 Glu Arg Leu Ala Lys Asn Leu Val Glu Lys Gly Val Leu Thr Thr Glu 115 Lys Gin Asn Phe Leu Leu Phe Asp Met Thr Thr His Pro Leu Thr Asn 110 158 150 Asp Lys Trp Val Asn Asp Pro His Arg Met Asp Arg Arg Leu Leu Ala 175 Leu Ile Tyr Leu Ala His Ala Ser Asp Val Leu Glu Asn Ala Phe Ala 180 Pro Leu Leu Asp Glu Gln Tyr Asp Leu Ala Thr Lys Arg Val Arg Gln 200 Leu Asp Leu Asp Pro Glu Val Glu Cys Leu Lys Ala Asn Thr Asn 210 Asn Asn Ile Lys Gln Arg Leu Ile Lys Lys Val Gln Glu Ala Val 145 ž ŝ Glu Val Leu Trp Ala Val Val Ala Ala Phe Thr Lys Xaa 225 (A) LENGTH: 200 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 488: Thr Gln Pro Pro Glu Thr Val Gln Asn Trp Ile Glu 85 90 (2) INFORMATION FOR SEQ ID NO: 488: 3

Met Ala Gln Arg Met Val 1rp Val Asp Leu Glu Met Thr Gly Leu Asp 1 $$\rm 10$ $\rm 15$ Ile Glu Lys Asp Gln Ile Ile Glu Met Ala Cys Leu Ile Thr Asp Ser 20

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Ser Met Ser Asp Trp Cys Lys Glu His His Gly 55 Lys Ala Val Lys Glu Ser Thr Ile Thr Leu Gln 70 75 80 Lys Ser Gly Leu Thr 65 Asp Glu Leu Leu Asp 50 55

Gln Ala Glu Tyr Glu Phe Leu Ser Phe Val Arg Gln Gln Thr Pro Pro

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Asp Leu Asn Ile Leu Ala Glu Gly Pro Asn Leu Ile Ile Lys Gln Pro

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Leu Asp Lys Tyr Met Pro Gln Phe Met Lys His Leu His Tyr Arg Ile 115 $$129\,$ Asp Ile Ser Glu Ser Ile Lys Glu Leu Gln Phe Tyr Arg Asn Asn Ile 170 175 Gly Leu Cys Pro Leu Ala Gly Asn Ser Val His Glu Asp Lys Lys Phe $100\,$ Ile Asp Val Ser Thr Val Lys Glu Leu Cys Arg Arg Trp Tyr Pro Glų 130 Glu Tyr Glu Phe Ala Pro Lys Lys Ala Ala Ser His Arg Ala Leu Asp 145 Phe Lys Lys Lys Ile Asp Glu Lys Lys Arg Lys Ile Ile Glu Asn Gly 180 으 2

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(2) INFORMATION FOR SEQ ID NO: 489:

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Glu Asn Glu Lys Thr Val Ser Xaa 195

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SEQUENCE DESCRIPTION: SEQ ID NO: 489 SEQUENCE CHARACTERISTICS:
(A) LENGTH: 351 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Ŧ 3 8

Gly Met Lys Gly Phe Lys Gly Gln Leu Ser Arg Gln Val Ala Asp Gln 89 90 95 Ala Asn 11e Asp 11e Leu Arg Pro Tyr Phe Asp Val Glu Pro Ala Gln 115 Val Arg Thr Gly Leu Leu Glu Ser Met Ile Pro Ile Lys Met Val Asn 130 Met Ala Thr Thr Ala Ala Pro Ala Gly Gly Ala Arg Aen Gly Ala Gly 1 5 10 15 Pro Glu Trp Gly Gly Phe Glu Glu Aan 11e Gln Gly Gly Gly Ser Ala $$20\ \ \ 25$ Val Ile Asp Met Glu Asn Met Asp Asp Thr Ser Gly Ser Ser Phe Glu 15 Asp Net Gly Glu Leu.His Gln Arg Leu Arg Glu Glu Glu Val Asp Ala 50 Asp Ala Ala Asp Ala Ala Ala Glu Glu Glu Asp Gly Glu Phe Leu 65 Leu Tyr Met Trp Gln Ala Gly Lys Arg Gln Ala Ser Arg Ala Phe Ser 100 35 6 S 55 45

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Phe Pro Gln Lys Ile Ala Gly Glu Leu Tyr Gly Pro Leu Met Leu Val 145 150 150

Phe

Thx Leu Val Ala Ile Leu Leu His Gly Met Lys Thx Ser Asp Thx $170\,$

Glu Ser Gly Thr Ser Gly Gly Gly Gly Ser Thr Glu Glu Ala Phe Met . 20 $$25\,$ Thr Ser Lys Asn Glq Ile Glu Arg Leu Thr Arg Pro Gly Ser Ser Tyr 50 55 Phe Tyr Ser Glu Val Lys Gln Ile Glu Lys Arg Asp Ser Val Leu 35 40 45

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Met Arg Gly Ser Arg Gly Gly Trp Ala Gly Glu Met Ala Ala Ser Gly
.1 5 10

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Ξ SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 490:

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Œ SEQUENCE DESCRIPTION: SEQ ID NO: 490: (A) LENGTH: 265 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Ala Thr Ala Lys Ala Val Ala Val Thr Leu Gln Ser His Xaa 340 345

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Asp Ile Pro Ala Met Leu Pro Ala Ala Arg Leu Pro Thr Thr Val Leu 325 330 335

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Asp Thr Leu Glu Gly Pro Asn Ile Pro Pro Ile Gln Arg Val Pro Arg 305 310

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Gly Leu Phe Gly His Cys Ile Val Leu Phe Ile Thr Tyr Asn Ile His 225 230 230

Leu His Ala Leu Phe Tyr Leu Phe Trp Leu Leu Val Gly Gly Leu Ser 245 250 255

Thr Leu Arg Met Val Ala Val Leu Val Ser Arg Thr Val Gly Pro Thr 260 260

Cys Asn Ala Gln Ile Thr Met Leu Gln Met Leu Ala Leu Leu Gly Tyr 210 225

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Ala Leu Asp Val Ile Gin Ala Gly Lys Glu Tyr Val Glu His Thr 130 140

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Val Glu Glu Asp Asp Pro Glu Leu 165

Phe Lys Gln Ala Val Tyr Lys Gln 170

Val Lys Glu Arg Lys Lys Gln Leu Lys Lys Glu Gly Lys Pro Thr Ile 145 150 150

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Pro Asp Lys Asn Gln Asp Asp Ala 100

Asp Arg Ala Gln Lys Ala Phe Glu 105 110

Asp Glu Glu Ile Lys Lys Arg Phe Arg Gln Leu Ser Ile Leu Val His 85 90 95

Phe Asn Leu Asn Pro Phe Glu Val Leu Gln Ile Asp Pro Glu Val Thr 65 70 75 80

Ala Val Asp Lys Ala Tyr Lys Leu Leu Leu Asp Gln Glu Gln Lys Lys 115 120 125

Gly Tyr Trp Leu Gly Val Ser Ser Phe Ile Tyr Phe Leu Ala Tyr Leu 200 205

Ile Ile Arg Glu Gly Thr Leu Met Gly Thr Ala Ile Gly Thr Cys Phe 180 180

5

Leu Leu Tyr Leu His Phe Ala Tyr His Lys Val Val Glu Gly Ile Leu $290 \hspace{1.5cm} 295 \hspace{1.5cm} 300 \hspace{1.5cm}$

Gln Arg Leu Leu Cys Gly Thr Leu Ala Ala Leu His Met Leu Phe 275 280 285

ઝ Glu Ser Arg Asp Gly Arg Val Asp Ser Trp. 225 230

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Ala Gln Glu Lys Ala Lys Arg Glu Arg Glu Trp Gln Lys Asn Phe Glu 210 225

Arg Asn Phe Gln Ala Asn 235 240

25

Met Lys

Leu Phe Ala Glu Leu Glu Ile Lys Arg Lys Glu Arg Glu 180 185 190

Lys

Glu Met His Glu Arg Lys Arg Gln Arg Glu Glu Glu Ile Glu 195 200

Thr Lys Gly Lys Lys Glu Lys Lys Asn Arg Thr Phe Leu Arg Pro Pro 255

2 INFORMATION FOR SEQ ID NO: 491: 6

Lys Val Lys Met Glu Gln Arg Glu Xaa 260 265

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 25 amino acids

25

Asp Ser Met Pro Thr Cys Pro Leu Xaa Ala Ser Leu Glu Cys Gly Pro 1 15 15 Ě SEQUENCE DESCRIPTION: SEQ ID NO: 491: (D) TOPOLOGY: linear (B) TYPE: amino acid

8

Leu Leu Pro Val Arg Leu Cys Cys Leu 20

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(2) INFORMATION FOR SEQ ID NO: 492:

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(A) LENGTH: 159 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 3

SEQUENCE DESCRIPTION: SEQ ID NO:

Asn Glu Tyr Arg Val Pro Glu Leu Asn Val Gln Asn Gly Val Leu 5 ş Ser Lys

9

Ser Phe Leu Phe Glu Tyr Ile Gly Glu Met Gly Lys Asp $20 \ \ \, 25$ Tyr Ile Tyr Ala Val Thr

Pro Leu Leu Glu Asp Ala Leu Met Asp Arg 40 Thr Ala Ser Ala Val Val Gin His Met Ser 55 60 Leu Val His Arg Gln 50 Asp 2

5 E

Gly Val Tyr Gly Phe Gly Cys Glu Asp Ser Leu Asn His Leu Leu 70 75 80 Asn Tyr Val Trp Pro Asn Val Phe Glu Thr Ser Pro His Val Ile Gln

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Ala Val Met Gly Ala Leu Glu Gly Leu Arg Val Ala Ile Gly Pro Cys 100

23

Met Leu Gln Tyr Cys Leu Gln Gly Leu Phe His Pro Ala Arg Lys 115 Arg

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Ser

Arg Asp Val.Tyr Trp Lys lie Tyr Asn Ser lie Tyr lie Gly 130 Leu ile Ala His Tyr Pro Arg ile Tyr Gln Arg Xaa 150 Gln Asp Ala I 145 35

INFORMATION FOR SEQ ID NO: 493: 3 6

(A) LENGTH: 279 amino acids TYPE: amino acid SEQUENCE DESCRIPTION: SEQ ID NO: 493: ž

45

Thr Ser Met Met Ile Ser Asp Asn Ser Ala Glu Asn Ile Ala Leu Val Tyr Asp Gly Leu Leu Gln Ala Gly Ala Arg Leu Cys Pro Thr Val Gln 20 30 20

Leu Glu Asp Ile Arg Asn Leu Gln Asp Leu Thr Pro Leu Lys Leu Ala 15 45 Ala Lys Glu Gly Lys Ile Glu Ile Phe Arg His Ile Leu Gln Arg Glu SO 60 55

Ser Gly Leu Ser His Leu Ser Arg Lys Phe Thr Glu Trp Cys Tyr 70 70 75 P 26

8

Gly Pro Val Arg Val Ser Leu Tyr Asp Leu Ala Ser Val Asp Ser Cys

Ser Pro Cys Lys Leu Glu Ile Ile Ala Phe His 105 Glu Glu Asn Ser Val S

His Arg His Arg Met Val Val Leu Glu Pro Leu Asn Lys Leu Leu Gln Leu Asn Phe Leu Cys Lys Trp Asp Leu Leu Ile Pro Lys Phe Phe Ala

0

Asn Leu lle Tyr Met Phe lle Phe Thr Ala Val Ala Tyr His Gln Pro 160 165 15

Leu Leu Thr Gly His Ile Leu Ile Leu Leu Gly Gly Ile Tyr $$180\$ Leu Lys Lys Gln Ala Ala Pro His Leu Lys Ala Glu Val Gly Asn 175 Met 뵱 Ser 2

Leu Leu Val Gly Gln Leu Trp Tyr Phe Trp Arg Arg His Val Phe Ile 195

Trp 11e Ser Phe 11e Asp Ser Tyr Phe Glu Ile Leu Phe Iln 210 215 Cys Phe Leu Xaa Ile Glu 235 Ala Leu Leu Thr Val Val Ser Gln Val Leu 235 22

Leu Asn 255 Leu Leu Tyr Thr Arg Gly Phe Gln His Thr Gly 11e Tyr Ser Val 260 Trp Tyr Leu Pro Leu Leu Val Ser Ala Leu Val Leu Gly Trp $245\,$ 2

Met Ile Gln Lys Pro Trp Xaa 275

33

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(2) INFORMATION FOR SEQ ID NO: 494:

(xi) SEQUENCE DESCRIPTION: SEQ.ID NO: 494: (A) LENGTH: 193 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 45

Met Ile Arg Cys Gly Leu Ala Cys Glu Arg Cys Arg Trp Ile Leu Pro 1 5 10 15 Arg Gly Trp Leu Gln Ser Ser Asp His Gly Gln Thr Ser Ser Leu Trp 35 Leu Leu Leu Ser Ala Ile Ala Phe Asp Ile Ile Ala Leu Ala Gly 20 20 55

Trp Lys Cys Ser Gln Glu Gly Gly Gly Ser Gly Ser Tyr Glu Glu Gly 8

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S 8

Cys Gln Ser Leu Met Glu Tyr Ala Trp Gly Arg Ala Ala Ala Ala Met 65 70 75 Leu Phe Cys Gly Phe Ile Ile Leu Val Ile Cys Phe Ile Leu Ser Phe 90 95 Phe Ala Leu Cys Gly Pro Gln Met Leu Val Phe Leu Arg Val Ile Gly 100 105 110

S

₹ Gly Leu Ala Leu Ala Ala Val Phe Gln Ile Ile Ser Leu Val Ile 115 120 125

Pro Val Lys Tyr Thr Gln Thr Phe Thr Leu His Ala Asn Xaa Ala 130 140

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Val Thr Tyr Ile Tyr Asn Trp Ala Tyr Gly Phe Gly Trp Ala Ala Thr 145 150

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Ile Ile Leu Ile Gly Cys Ala Phe Phe Phe Cys Cys Leu Pro Asn Tyr 165 170 175

Glu Asp Asp Leu Leu Gly Asn Ala Lys Pro Arg Tyr Phe Tyr Thr Ser 180

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Met Ala Leu Thr Leu Leu Pro Ser Val Ser Arg Leu Pro Gly Glu Arg

35

Met Lys Val Ile Phe Phe Pro Tyr Pro Val Leu Pro Leu Pro Ala Pro 35 $40\,$ 45

Met Ala Ala Ser Gly Lou Pro Tyr Val Lou His His Lys Ser Ser Leu
20 25 30

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SEQUENCE DESCRIPTION: SEQ ID NO: 496:

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(2) INFORMATION FOR SEQ ID NO: 496:

SEQUENCE CHARACTERISTICS

(A) LENGTH: 147 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

5

Arg Lys Gln Ala Ser Pro His Arg Ile Leu Phe His Xaa 195 200 205

5

Pro Phe Ser Pro Pro Ala Cys His Thr Ala Pro Asn Ser Val Leu Ile 145 150 150

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Pro Pro Pro Asp Ser Asp Leu Cys Ser Gly Pro Leu Leu Pro Gly
130 140

Gln Ser Leu Phe Cys Lys Ser Glu Leu Trp Trp Arg Gln Met Arg Ser 165 170 175

Ile Thr Trp Val Pro Ser Pro Lys Ala Gly Trp Arg Trp Thr Lys Gly 185 190

(2) INFORMATION FOR SEQ ID NO: 495:

Ξ SEQUENCE CHARACTERISTICS:

(A) LENGTH: 205 amino acids(B) TYPE: amino acid

Met Ala Ala Gly Asp Gln Val Phe Ser Gly Ala Gly His Val Xaa Glu 1 15

8

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Gly Thr Trp Val Pro Arg Leu Val Leu Gly Leu Gly Ser Gly Asp $50 \ \ \, 55$

Gln Val His Tyr Leu Pro Ile Ser Ser Ser Ile Val Asn Tyr Gly Thr $65 \qquad 70 \qquad 75 \qquad 80$

45

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Pro Thr

Trp Ser Thr Arg Cys Phe Gln Val Trp Asp Leu Leu Ser 100 105

Ser Val Ser Gly Lys Ser Trp Val 85

Phe

Leu Val Tyr Pro Leu His Pro 90 95

His Val Ala Gly Gly Arg His Ala Trp Leu Leu Thr Trp Gln Ser Ala 25 30

Cys Pro Ala As
n Arg Leu Ser Leu Val Pro Leu Val Pro Ser Ala Ser 35 $40\,$

Met Thr Arg Leu Met Arg Xaa Arg Thr Ala Ser Gly Ser Ser Val Ile 50 55

Leu Trp Met Ala Pro Ala Ala Ala Pro Thr 65 70 Pro Ala Arg Ala Pro Glu 75

Ala Ala Pro Thr Pro Ala Arg Ala Pro Ala Ala Ala Arg Thr Pro Ala 95

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SEQUENCE DESCRIPTION: SEQ ID NO: 495: (D) TOPOLOGY: linear

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8 Arg Gly Pro Thr Trp Thr Ser Pro Pro Thr Arg Val Leu Leu Gly Thr 105 100 1 105

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Va.

Pro Gly Leu Ser Gln Leu Pro Thr Ser His Lys Pro Ile Lys Gln 130 140

Glu Tyr Xaa 145

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Glu Leu Pro 115

Asp Lys

Gly Glu Gly Asn Thr Arg Arg Ala Ser Gly 120 125

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Xaa Pro Gly Pro Ser Pro Trp Arg Ser Pro Ala Arg Arg Pro Ala Gln
115 120 125

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(2) INFORMATION FOR SEQ ID NO: 497:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 64 amino acids (B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 497:

Met Val Trp Val Leu Trp Ser Ala Pro Ser Leu Ala Pro Pro Trp Val 1 5

9

Gly Pro Cys Trp Pro Ser Thr Gly Asn Cys Cys Leu Cys Glu Val Gly 20 25

Ala Ala Leu Pro Pro Arg Gly Pro Ser Leu Ser Asp Cys Leu Gly. Leu 15

Pro Pro Trp Thr Pro Trp Gly Pro Ala Trp Thr Leu Ala Gln Ser Xaa 50 60 2

22

(2) INPORMATION FOR SEQ ID NO: 498:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 94 amino acids

3

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 498: (B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Ser Thr Gly Ala Leu Asn Thr Ser Pro Pro Ala Ser Asn Arg Leu 35 Glu Ser Thr Leu Asn Glu Tyr Leu Ile Gln Pro Gln Leu His Cys Ser

Ser Val Gln Arg Leu Thr Leu Lys Trp Gly Cys Ser Ser Leu Gln Arg 45 6

Asp Gly Gln Ala Val Pro Trp Gly Leu Trp Gln Arg Ala Tyr Pro Ser 50 60

Pro Ser Asp Leu Leu Arg Pro His Ala Val Thr 70 80 星 Leu Leu Pro Thr

5

Xaa Ser Val His Thr Cys Glu Ser Ser Pro Ser Val Ser Val 20

(2) INFORMATION FOR SEQ ID NO: 499: 55

(A) LENGTH: 22 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: 1inear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 499:

8

Met Phe Leu Ile Phe Val Tyr Phe Leu Lys Xaa Leu Phe Ser Ser Ser 15

Leu Pro Phe Leu Trp Leu 20

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(2) INFORMATION FOR SEQ ID NO: 500: 2

(A) LENGTH: 33 amino acids (i) SEQUENCE CHARACTERISTICS:

TYPE: amino acid (D) TOPOLOGY: linear

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 500:

Arg Gly Gly Leu Cys Pro Leu Leu Val Pro Gly Pro Leu Ala Arg Gln

Glu Pro Ser Pro Ser Leu Gln Gly Cys Ser Glu Ser Pro Val Gly Met 8

2

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52

(2) INFORMATION FOR SEQ ID NO: 501:

(i) SEQUENCE CHARACTERISTICS:

9

(A) LENGTH: 28 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 501:

35

Met Gln Phe Leu Leu Thr Ala Phe Leu Leu Val Pro Leu Leu Ala Leu

Cys Asp Val Pro Ile Ser Leu Gly Phe Ser Pro Ser 20 25

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(2) INFORMATION FOR SEQ ID NO: 502:

45

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 15 amino acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 502: (X

Pro Gly Lys Pro Gln Ala Cys Pro Glu Leu Thr Ser Val Leu Pro 1 10 11

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(2) INFORMATION FOR SEQ ID NO: 503:

(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 19 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Ž. SEQUENCE DESCRIPTION: SEQ ID NO: 503:

S

Asn Lys Ser Leu Xaa Ser Cys Leu Phe Val Leu His Phe Val Leu His 1 15

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Leu Gly Pro Arg Gln Leu Pro Glu Pro Ser Gln Gln Ser Lys Xaa 195 200 205

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Ala Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly
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Pro Ser Ser Phe Gln Asn Pro Ala Ser Ser Pro Ser Ser Trp Thr His 50 . 60

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Asn Leu Ser

Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp Thr Leu 20 25 30

Met Val Asp Ile Ser Lys Met His Met Ile Leu Tyr Asp Leu Gin Gin 1 . 15

40

Ala Phe Arg Gln Val Arg Leu Lys His Arg Lys Leu Arg Glu Gln Val 130 135

Ser His Ala Ala Arg Arg His Gln Arg Xaa Leu Leu Ala Ala Ile Asn 115 120 125

\$

Gln Gln Asn Leu Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp 165 170 175

Thr Leu Ala Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala 180

Asn Ser Met Val Asp Ile Ser Lys Met His Met Ile Leu Tyr Asp Leu 145 150 150

35

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SEQUENCE DESCRIPTION: SEQ ID NO: 505:

30

(2) INFORMATION FOR SEQ ID NO: 505:

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Met Met Asp Ile Gln Tyr Thr Lys Glu Met Lys Glu Ser Ala Ala Arg 85 90 95

25

Thr Gly Val Met Gly Val Cys Cys Thr Ala Leu Leu Val Ala Val Val 50 55 60

Ala Arg Lys Leu Glu Phe Asn Lys Ala Glu Lys His Val His Asn Phe 65 70 75 80

Gly Asp Val Val Pro Gly Thr Met Trp Gly Lys Ile Val Cys Leu Cys 35 40 45

Ser Asp Thr Leu Trp Leu Ile Pro Ile Thr Phe Leu Thr Ile Gly Tyr $20 \ 25 \ 30$

35

VAL

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Gln Glu Ala Trp Met Phe Tyr Lys His Thr Arg Arg Lys Glu 100 105 110

(i) SEQUENCE CHARACTERISTICS:

(a) LENGTH: 75 amino acids
(b) TYPE: amino acid (D) TOPOLOGY: linear

25

Ser Ile Leu Asn Leu Phe Leu Leu Lys Met Ile Val Ser 20

Met Glu Lys Thr His Arg Leu Arg Ile Arg Asn Pro Cys Leu Gln Phe 1 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 504:

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15

Met Leu Trp Phe Gly Gly Cys Ser Ala Val Asn Ala Thr Gly His Leu 1 5 10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 507:

(A) LENGTH: 207 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

5

(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 507:

5

(2) INFORMATION FOR SEQ ID NO: 504:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 29 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Cys Xaa Phe

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INFORMATION FOR SEQ ID NO: 508:

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(2) INFORMATION FOR SEQ ID NO: 506:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 10 amino acids
(B) TYPE; amino acid
(D) TOPOLOGY: linear

50

Glu Glu Glu Pro Gly Tyr Phe Pro Gln Tyr Xaa 65 70 75

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 36 amino acids

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Leu Pro Leu Ala Glu Leu Lys Asn Trp Val 1 5 10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 506:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 508: (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Trp Arg Cys Arg Gly Lys Leu Ser Phe Pro Leu Phe Ala Val Val 1 5 10 15 ile Val Ser Cys Arg Lys Asp Gly Pro Asp Ala Ala Ala Ala Pro Ala 25 2

Val Xaa Lys Lys 35

(2) INFORMATION FOR SEQ ID NO: 509: 5

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 19 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 509: (B) TYPE: amino acid (D) TOPOLOGY: linear 8

Met Ala Leu Val Ala Leu Phe Thr Gln Leu Met Arg Xaa Leu Gly Arg 25

Cys Pro Gln

3

(2) INFORMATION FOR SEQ ID NO: 510:

(A) LENGTH: 32 amino acids (1) SEQUENCE CHARACTERISTICS: 35

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 510:

Met Thr Phe Pro Phe Glu Lys Glu Asn Ser Cys Phe Gln Cys Leu Leu 1 1 15 $\,$

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Phe Asp Ser Trp Arg Glu Gln Thr Arg Thr Asn Ile Gln Pro Gln Arg

5

(2) INFORMATION FOR SEQ ID NO: 511:

20

(A) LENGTH: 28 amino acids SEQUENCE CHARACTERISTICS:

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 511: (B) TVPE: amino acid (D) TOPOLOGY: linear

Met His Leu Leu Asp Phe Phe Arg Asp Leu Val Leu Leu Val Leu 115 $^{\rm 15}$

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299

Ala Leu Leu Asp Ser Phe Trp Leu Glu Val Gln Lys 20

(2) INFORMATION FOR SEQ ID NO: 512:

(A) LENGTH: 26 amino acids (i) SEQUENCE CHARACTERISTICS:

2

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 512: (D) TOPOLOGY: linear

Met Cys Leu Ile His Phe Ile Lys Ile Ile Leu Val Phe Ile Leu Lys

12

Leu Trp Leu Tyr Ser Gln Lys Cys Pro Lys 20 25

2

(2) INFORMATION FOR SEQ ID NO: 513:

(A) LENGTH: 33 amino acids (i) SEQUENCE CHARACTERISTICS:

23

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 513:

30

Met Ile His Val His Glu Trp Asn Asp Gln Met Leu Met Val Tyr Ile

Phe Leu Tyr Pro Val Ser Ile Thr Phe Leu Asn Leu Cys Ser Leu Thr 35

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(2) INFORMATION FOR SEQ ID NO: 514:

(A) LENGTH: 47 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

45

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 514:

Leu Asn Glu Ser Tyr Val Ser Arg Ala Gly Gly Try Phe Ser Met Phe

S

Xaa Leu Ile Phe Phe Leu Leu Ala Leu Gly Ser Xaa Leu Cys Leu Leu 20

25

Leu Cys Leu Pro Ser Phe Asn Lys Thr Arg Arg Lys Gln Lys Pro

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(2) INFORMATION FOR SEQ ID NO: 515: (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids

(ž SEQUENCE DESCRIPTION: SEQ ID NO: 515: (B) TYPE: amino acid
(D) TOPOLOGY: linear

S

Ser Ser Lys Thr Pro Leu Pro Ser Glu Arg Arg Trp Ile Ser Gly Ser 1 10 15

5

Val Pro Pro Asn Ala Pro Pro Thr Leu Thr Ile Lys Leu Leu
165 170

Thr Trp Ala Pro Ser Ala Ala Ala Ser Cys Ala Cys Ile Met Thr Glu 145 150 150

S

Ser

Val His Ser Leu Asp Gly Phe Arg His Gln Ala Ser Gly Thr Ala 130 140

5

reu Met Ala Pro Arg Pro Trp Leu Leu Gly Ile Ala Leu Leu Gly 20 $25\,$ 30

Leu į Ala Leu Glu Pro Ala Leu Gly His Trp 35 40

7

(2) INFORMATION FOR SEQ ID NO: 516: (i) SEQUENCE CHARACTERISTICS:

8

(A) LENGTH: 3 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

25

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SEQUENCE DESCRIPTION: SEQ ID NO: 516:

25

Met Trp Lys Asn Leu Gly Ser Gly Ser Val Phe Val Thr Trp Phe Ser 1 10 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 518:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 43 amino acids

Leu Val Met Ile Leu Ser Gly Ile Gly Pro Leu Gly Asp Ala Glu Asp 20 25

20

5

(2) INFORMATION FOR SEQ ID NO: 518:

(i) SEQUENCE CHARACTERISTICS:

30 Ser Ile Ser Asp Val Ser His Arg Leu Arg Pro 35 40

35 (2) INFORMATION FOR SEQ ID NO: 519:

9 SEQUENCE CHARACTERISTICS:

(A) LENGTH: 13 amino acids

Ě SEQUENCE DESCRIPTION: SEQ ID NO: 519: (B) TYPE: amino acid
(D) TOPOLOGY: linear

6

Phe Gln Phe Pro Leu Leu Thr Ile Ala Leu Gln Phe Leu 1 5 10

3

(2) INFORMATION FOR SEQ ID NO: 520

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Val Asp Ser Cln Met Asp Asp Met Asp Met Asp Leu Asp Lys Glu Phe 20 30

Phe Ala Phe Cys Ala Glu Leu Met Ile Gln Asn Trp Thr Leu Gly Ala 1 10 15

SEQUENCE DESCRIPTION: SEQ ID NO: 517:

(D) TOPOLOGY: linear

55

S

Leu Phe Val Asp Leu Val Glu Lys Phe Val Glu Pro Cys Arg Ser Asp 100

Leu Val Asn Val Ala Ala Lys Leu Thr His Asn Lys Asp Val Arg Asp 95

Gly Val Phe Ser Glu Met Glu Ala Asn Phe Lys Asn Leu Ser Arg Gly 65 70 75 80

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Leu Asp Leu His Lys Ser Leu Val Cys Thr Ala Leu Arg Gly Lys Leu 50 55 60

Leu Gln Asp Leu Lys Glu Leu Lys Val Leu Val Ala Asp Lys Asp Leu 35 $40\,$

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 174 amino acids (B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 517:

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Leu Asn Trp 1

(1) SEQUENCE CHARACTERISTICS:

8

(A) LENGTH: 30 amino acids(B) TYPB: amino acid(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 520:

55 Met His Tyr Val Ile Val Leu Ser Leu Phe Val Val Leu Glu Lys Lys
1 10 15

8 Asn Lys Met Gly Ser Asp Gly Cys Leu Arg Lys Asn Gly Ser 20 25 30

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His Trp Pro Leu Ser Asp Val Arg Phe Phe Leu Asn Gln Tyr Ser Ala 115 120 125

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Asn Arg Thr Leu Leu Phe Leu Ile Leu Phe Val Leu Phe Gly Leu Gly

Tyr Gly Phe

15

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(2) INFORMATION FOR SEQ ID NO: 525:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 524:

(A) LENGTH: 19 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 524:

Met Ser Arg Ser Ile Val Leu Arg Gly Ser Leu Phe Leu Phe Ser His Tyr Thr Leu Lys Leu Leu Ser Val Ile Lys Gln Thr Asn Arg Lys Pro Leu Pro Val Leu Leu Cys Leu Thr Leu Pro Met Pro Leu Pro 15 Leu Leu Ile Thr Glu Ser Gly Ser His Glu Lys Lys Ser Phe Tyr Pro 20 Lys Tyr Met Phe Lys Ile Ile Ile Tyr Val Ser Ala Tyr Cys 35 Ser Ser Ile Pro Val Ser Ile Leu Ile Gly Met Lys Leu Ile Leu Tyr Ile Val Trp Glu Lys Pro Cys Ile Arg Leu Phe Tyr Xaa Val Leu 35 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 521: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 522; (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 523: 965 (A) LENGTH: 47 amino acids (A) LENGTH: 26 amino acids (A) LENGTH: 58 amino acids Ser Ala Thr Ala Arg Gly Gly Asn Arg Thr 20 25 (i) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (B) TYPE: amino acid (2) INPORMATION FOR SEQ ID NO: 523: (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 521. (D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 522. (D) TOPOLOGY: linear (D) TOPOLOGY: Linear WO 98/39448 3 Ser Phe Met 2 15 2 23 3 35 S 45

Phe Leu Leu Val Leu Ser Val Phe Cys Asp Phe Met Cys Ser Ile 5 $$\rm 10$ Ala Pro Arg Cys His Ala Leu Ser Leu Val Ser Leu Arg Ala Gln His 20 Val Ser Ser Trp Lys Thr Phe Leu Pro His Phe Ser Leu Pro Gly Pro 25 20 30 Met Leu Leu Phe Ile Leu Leu Thr Leu Ser Ser Gly Cys Arg Leu Leu 1 5 Arg Glu His Pro Glu Gly Ser Arg Thr Trp Phe Phe Arg Tyr Trp Glu 35 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 525: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 526: (A) LENGTH: 40 amino acids (A) LENGTH: 57 amino acids (i) SEQUENCE CHARACTERISTICS: (1) SEQUENCE CHARACTERISTICS: Pro Gly Ala His Cys Leu His Cys Ala 50 55 (B) TYPE: amino acid (D) TOPOLOGY: linear (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 526 Leu Phe Ile Thr Cys His Ser Met Leu 2 52 3 6 5 35 S 55

(1) SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 527:

Arg Thr Ala Leu Arg Ala Thr Val Ser His

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5 Ser Cys Asn Gln Leu 20 Ala Arg Leu Leu Phe Leu Ser Ser Val His Pro Ser Ile Met Pro 1 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 527: (A) LENGTH: 21 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

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Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 39 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 528:

20

Pro Gln Lys Leu Gln Phe Leu Ser Trp Met Glu Arg Gln Gln Arg Cys \$25\$

Thr Gly Val Ala Lys Tyr Ala

30

(2) INFORMATION FOR SEQ ID NO: 529:

Ξ SEQUENCE CHARACTERISTICS:

35

(<u>x</u> SEQUENCE DESCRIPTION: SEQ ID NO: 529:

(A) LENGTH: 128 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

6

Met Val Leu Arg Leu Ile Gln Leu Ile Phe Leu Ile Phe Phe Ile His 1 5 10

Ile Ile Ile Leu Leu Ile Pro Gly Ser Arg Pro Cys Gly Ser Trp Val 20 25

2

Asn Asp Arg Xaa Leu Gly Leu Arg Asp Val Thr His Leu Ile Tyr Leu 35 $40\,$

His Trp Val His Gly His Leu Pro Trp Cys His Pro Tyr Ile Gln Val 50 55

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Glu Phe Ser Ala Leu Ile Glu Ser Thr Ala Gln Leu Gly Leu Pro Phe 65 70 75 80

Ser Trp Val Arg Val Ile His Pro Phe Leu Val Leu Pro Cys Leu Tyr 85 90 95

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Ser Pro Gly Leu Lys Asn(Gly Ile Phe Leu Phe Leu Leu Arg Ala Met $100\,$

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(2) INFORMATION FOR SEQ ID NO: 528:

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X X

Met Ser Leu Thr Ser Ser Leu Thr Phe Leu Ser His Ile Leu Leu Leu 1 5 10

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30 Pro Pro Asp Gln Glu Pro Gln Lys Glu Thr Glu Pro Ala Thr Ser Ser 50 55 25

Trp Gly Phe Gly Xam Lys Thr Alm Ser Phe Gly Alm Val Gly Glu Thr
35 40 45

Leu Gly Gly Arg Cys Ile Gln Gly Arg Phe Ala Ser His Ser Lys Phe
20 25 30

20

Met Gly Ser Ser Val Leu Pro Phe Cys Val Cys Val Thr Ser Pro Ser

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SEQUENCE DESCRIPTION: SEQ ID NO: 530: (A) LEWSTH: 82 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear 0

(2) INFORMATION FOR SEQ ID NO: 530:

(i) SEQUENCE CHARACTERISTICS:

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His Ala Arg Pro Trp Ala Arg Val Ile Gly Leu Arg Ile Trp Pro Oln 65 70 75

Pro Asn

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(2) INFORMATION FOR SEQ ID NO: 531:

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Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 20 amino acids

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 531:

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Met Leu Leu Ser Val Ala Ile Phe Ile Leu Leu Thr Leu Val Tyr Ala 1 5 10 15

Tyr Trp Thr Met 20

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(2) INFORMATION FOR SEQ ID NO: 532:

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(1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 75 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 532: (D) TOPOLOGY: linear

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Pro Gly Gly Met Phe Pro Gly Asn Leu Glu Ala Phe Arg Val Pro Val 115 120 120

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Asn Cys Glu Ile Leu Glu Tyr Cys Tyr Tyr Leu Thr Gln Leu Lys Ile Arg Cys Ser Ile Thr Ala Val Ser Asp Ser Ser Thr Ser Trp Ala Ile Lys Ala Gln Leu Lys Ile Glu Asn Lys Asp Leu Asp Asn Lys Thr Ala 50 Ser Met Gly Lys Tyr Leu Ser Ile Pro Thr Val Leu Leu Lys Ile Ile 9

(2) INFORMATION FOR SEQ ID NO: 533: ឧ

Lys Gly Gly Gly Gln Glu Ala Leu Thr Cys Thr 65 75

15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 533: (A) LENGTH: 60 amino acids (1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 23 Met Phe Leu Met Arg Met His Leu Cys Phe Cys Lys Tyr Cys Cys Ser 1 1 Phe Ile Val Thr Pro Thr Ser Thr Ser Asn Thr Xaa Ser Tyr Leu Trp 20 3

Pro Trp Ile Ser Ala Ser Met Ala Gly Arg Gly Ser Xaa Trp Ala Cys Thr Leu Asn Ala Val Thr Arg Glu Gly Leu Pro Glu 50 50 35

(2) INFORMATION FOR SEQ ID NO: 534: 6

(A) LENGTH: 39 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid

45

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 534: (D) TOPOLOGY: linear

Ser Leu Leu Asn Thr His Thr Leu Cys Phe Val Leu Phe Cys Phe 5 The Leu Ser Ile Asn Gln Glu Lys Leu Ala Asn His Leu Ala Phe Arg 20 Mat S

lle Leu Phe Phe Ile Val Phe 35 55

(2) INPORMATION FOR SEQ ID NO: 535: 9

(A) LENGTH: 2 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 535: (D) TOPOLOGY: linear

Met Leu

9

(2) INFORMATION FOR SEQ ID NO: 536:

15

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 36 amino acids
(B) TYPE: amino acid

(b) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 536:

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Met Asp Gln Phe Lys Ile Phe Tyr Phe Leu Lys Ala Phe Phe Ala Cys 1 5 Cys Asn Val Gln Asp Pro Ser Pro Phe Met Gly Glu Thr Gly Ser Tyr 20 20

Leu Asn Ile Gly

22

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(2) INFORMATION FOR SEQ ID NO: 537:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 537: (A) LENGIH: 14 amino acids (1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 35

Met Phe Asp Phe Leu Ser Tyr Phe Lys Asp Leu Leu Ser Cys $_1$

6

(2) INFORMATION FOR SEQ ID NO: 538: 45

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 amino acids TYPE: amino acid (D) TOPOLOGY: linear

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Met Gly Phe Gly Phe Val Leu Asn Ile Phe Ser Phe Phe Leu Xaa Pro 12 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 538:

Pro Leu

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(2) INFORMATION FOR SEQ ID NO: 539:

55 છ 20 15 5 8 50 6 ઝ 25 3 Leu Leu Tep Thr Leu Leu Ala Xaa Tyr Xaa 1 5 10 Phe Val Leu Gln Arg Leu Val Arg Asn Val Glu Tyr Tyr Gln Ser Asn Tyr Val Phe 65 70 75 Glu Typ Leu Glu Axg Arg Arg Ala Thr Ile Axg Pro Typ Ser Thr Phe \$35\$Met Ala Ala Gin Lys Asp Gin Gin Lys Asp Ala Giu Ala Giu Gly Leu 1 15 2 Val Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met Leu 95 reu (2) INFORMATION FOR SEQ ID NO: 541: Leu Arg Thr Leu Glu Ser Lys Leu Val Leu Phe Gly Arg Glu Val Ser 35 40 INFORMATION FOR SEQ ID NO: 540: Val Ala Leu Ala Val Phe Phe Gly Ala Cys Xaa 100 105 Leu Val (i) SEQUENCE CHARACTERISTICS: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 541: . E Ě Ξ ž Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met $10 \ \, 15$ SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: 539: SEQUENCE DESCRIPTION: SEQ ID NO: 540: Ala Leu Ala Val Phe Phe Gly Ala Cys Tyr Ile Leu Tyr 20 25 (A) LENGTH: 11 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 108 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 106 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

5 S Pro Ala His Gln Tyr Ala Leu Ala Gly Gly Ile Ser Phe Pro Phe Phe 50 $\,$ 50 $\,$ Asp Gly Glu Glu Leu Gln Met Glu Pro Val 100 105 Leu Val Val Ile Gly Ser His Ala Ala Phe His Gln Ile Glu Ala Val 95 95 Trp Leu Ala Gly Ala Gly Ser Ala Val Phe Trp Val Leu Gly Ala Thr $65 \qquad 70 \qquad 75 \qquad 80$

5 (2) INFORMATION FOR SEQ ID NO: 542:

ઝ 30 25 20 Val Thr Leu Gly Ser Ser Phe Ile Pro Gly Glu His Phe Trp Leu Leu 65 70 75 80 Tyr Asn Arg Lys Tyr Leu Met Cys Gly Gly Ile Ala Phe Trp Ser Leu 50 60 Ser Ser Phe Asn Ile Gly Asp Ser Ser Ser Gly Leu Ile Gln Thr Val Phe Ile 20 25 Met Asp Arg Phe Thr Val Ala Gly Val Leu Pro Asp Ile Glu Gln Phe 1 15 (i) SEQUENCE CHARACTERISTICS Tyr Met Val Leu Ala Pro Val Phe Gly Tyr Leu Gly Asp Arg 35 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 542: (A) LENGTH: 136 amino acids
(B) TYPE: amino acid

Ala Thr Leu Gln Ala Pro Lys Xaa 130 135 Gly Cys Ser Ala Ser Ser Thr Leu Pro Phe Arg Trp Ala Val Val Trp 115 120 120

25

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Leu Leu Thr Arg Gly Leu Val Gly Val Gly Glu Ala Ser Tyr Ser Thr 85 90 95

Ile Ala Pro Thr Leu Ile Ala Asp Leu Phe Val Ala Asp Gln Arg Thr 100 105

5 INFORMATION FOR SEQ ID NO: 543: 50

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(1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 543: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 424 amino acids

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Met Ala Gly Asp Trp His Trp Ala Leu Arg Val Thr Pro Gly Leu Gly 1 10 15

Leu Val Val Arg Glu Pro Pro Arg 25

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Val Val Ala Val

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Ser

Leu Asn Pro Thr

Ser Asp Leu Pro Pro 40

Ala Val Glu Arg His 35

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Pro Leu Thr Tyr Leu His lle Cys His Ser Xaa Pro Trp Ala His Pro 335

Thr Lys Gly Leu Gly Leu Thr Pro Trp Pro Gly Pro Ala Ser Arg Gly 345

Ser Gly Ser Thr Leu Gly Arg Val Pro Ala Pro Arg His Tyr Xaa Gly 365 365 Glu Glu Val Gly Val Gln Glu Gly Asp Pro Ser Pro Gln Gly Xaa Pro 370 2

Gin Val Pro Ala Leu Val Phe Leu Try Val Ala Ser Asp Leu Ala Pro 415 Gin Giy Leu Giy Ala Ile Cys Asn Giy Ile Lys Phe Val Ala Arg Pro 385

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Leu Ala

Thr Ale Val Ala Phe Val Thr Gly Ser

Phe

Leu Gly

Ser

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Trp Ala Asp Leu Arg Ala Leu Ala Arg Asn Pro Ser Phe Val Leu 50

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Ser Arg Val Val Leu Gly Glu 90 95

Phe Leu Leu Arg

Leu Trp Ala Pro Ala 85

Ser Leu

Thr Pro Pro Cys Leu Pro Gly Asp Ser Cys Ser Ser Asp 100

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Leu His Pro Arg Ala Pro Glu P.F.B

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(2) INFORMATION FOR SEQ ID NO: 544;

(A) LENTH: 39 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

SEQUENCE DESCRIPTION: SEQ ID NO: 544: (X

Phe Arg Phe Val Ile Cys Leu Phe Leu Trp Leu Val Leu Cys Arg 10 Met

Ser Ala Ser Arg Ile Ala Leu Tyr Tyr Arg Ile Val Phe 20 Ser Thr βsp 35

(2) INFORMATION FOR SEQ'ID NO: 545:

Met

Tyr Leu Leu Phe Val Gln Phe Ser Pro Ala Phe Ser Arg Thr Ser Pro 20 55

Ser Gly Cys Ala Ala Arg Trp Leu Phe Ser

ile Phe Gly Leu Ile Thr Cys Leu Thr Gly Val Leu Gly Val Gly Leu 115

Gly Val Glu Ile Ser Arg Arg Xaa Arg His Ser Asn Pro Arg Ala Asp 130

22

Thr Gly Leu Leu Gly Ser Ala Pro Phe Leu Phe 150

Leu Val Cys Ala

Pro 145

3

Ser Ile Val Ala Thr Tyr Ile Phe 170

Leu Ala Cys Ala Arg Gly

Leu Ser

Ser Met Asn Trp Ala Ile Val Ala 185

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ile Phe Ile Gly Glu Thr Leu 180

35

Leu Tyr Val Val Ile Pro Thr Arg Arg Ser Thr Ala Glu 205

Leu 195

Asp 11e

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Ala Phe Gin Ile Val Leu Ser His Leu Leu Gly Asp Ala Gly Ser Pro 210

25

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Ser Leu Ile His Gln Cys Ser 35

8

(A) LENGTH: 58 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

45

Pro Pro 240

Tyr Leu Ile Gly Leu Ile Ser Asp Arg Leu Arg Arg Asn Trp 225

Leu Ser Glu Phe Arg Ala Leu Gln Phe Ser Leu Met Leu Cys 245 25

Ser Phe

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Gly Ala Leu Gly Gly Ala Leu Ser Trp Ala Pro Xaa Ser 260

Ala Phe Val

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Lys Gln Gly Pro Gln Thr Thr Gly Leu Trp Cys Pro Sar Gly 295 300

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Ser Leu Arg Pro Thr Ala Gly Gly His Ser Cys Thr Cys Arg Ala Cys 275

Ala Ala Pro Pro Ala Cys Pro Trp Pro Val Cys Ser Ser Glu Ary Leu 315

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 545:

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Leu Pro Trp Xea Ala Gin Leu Leu Asp Arg Thr Ile Gly Pro Leu 5

Trp Arg Ser Pro Lys Asn Phe Arg Arg Leu Tyr Pro Pro Cys Thr Thr 35

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(2) INFORMATION FOR SEQ ID NO: 546:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 33 amino acids

(B) TYPE: amino acid

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Œ SEQUENCE DESCRIPTION: SEQ ID NO: 546:

Met Gly Leu Ser Val Leu Leu Pro Leu Cys Leu Leu Gly Pro Gly Arg 1 10 15

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 547:

phe Thr Ser Gly Gln Lys Pro Leu Agp Thr Pro Gly Leu Gly Val Pro 20 30

8 Val Leu Asp Ala Asn Cys Ser Arg Asp Val Lys Gln Met Leu Leu Lys 245 250 255

5

5

Lys Leu Leu Lys Leu Thr Gly Ser Val Leu Glu Asp Ala Trp Lys Glu 210 225

Lys Gly Lys Met Asp Met Glu Glu Ile Ile Gln Arg Ile Glu Asn Vel 225 230 230

Asn Ala Leu Phe Ser Asn Pro Met Asp Asp Asn Leu Ile Cys Ala Val . 200 205

Gin Val Thr Arg Ala Asp Ile Leu Gin Val Gly Leu Arg Giu Leu Leu 180

Phe Leu Gly Glu Leu Tyr Leu Asn Leu Glu Ile Lys Gly Thr Asn Gly
175

Leu Val Glu Leu Arg Ser Ser Asn Trp Gly Arg Val His Ala Thr Ser 260 265 270

25 Thr Tyr Arg Glu Ala Thr Pro Glu Asn Asp Pro Asn Tyr Phe Met Asn 275 280 285

Glu Pro Thr Phe Tyr Thr Ser Asp Gly Val Pro Phe Thr Ala Ala Asp 290 300

ઝ Pro Asp Tyr Gln Glu Lys Tyr Gln Glu Leu Leu Glu Arg Glu Asp Phe 305 310 315 Phe Pro Asp Tyr Glu Glu Asn Gly Thr 325 Asp Leu Ser Gly Ala Gly Asp 330 335

33 Pro Tyr Leu Asp Asp Ile Asp Asp Glu Met Asp Pro Glu Ile Glu Glu 340 345 350

6 Ala Tyz Glu Lys Phe Cys Leu Glu Ser Glu Arg Lys Arg Lys Gln 355 360

(2) INFORMATION FOR SEQ ID NO: 548:

E SEQUENCE CHARACTERISTICS

45

(A) LENGTH: 77 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 548:

Met Leu Arg Leu Asp Ile Ile Asn Ser Leu Val Thr Thr Val 1 5 Phe Met

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Leu Ile Val Ser Val Leu Ala Leu Ile Pro Glu Thr Thr Thr Leu Thr $20 \ 25 \ 30$

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Leu Leu Gln Arg Cys Arg Thr Glu Tyr Glu Val Lys Asp Gln Ala 130 135

Leu Ser His His Leu Thr Ile Ser Pro Gln Ser Gly Asn Phe Arg Gln 115 120 125

Ala Lys Gly Asp Glu Val'Thr Arg Lys Arg Phe His Ala Phe Val Leu 145 150 150

8

S

Thr Ser Ile Pro Asn Phe Ser Tyr Met Gly Ala Arg Leu Cys Asn Tyr 100 105

3

Glu Thr Glu Ile Glu Gln Phe Ala Glu Thr Leu Asn Gly Cys Val Thr .65 $70\,$ $75\,$

The Asp Asp Ala Leu Gin Glu Leu Val Glu Leu Ile Tyr Gin Gin Ala 95

Tyr Val Gln Asp Phe Leu Asn His Leu Thr Glu Gln Pro Gly Ser Phe 50 60

Thr Glu Ser Tyr Glu Asp Gly Cys Glu Asp Tyr Pro Thr Leu Ser Glu 35

Ser Val Asn Ala Pro Glu Phe Tyr Pro Ser Gly Tyr Ser Ser Ser Tyr 20 30

Met Ala Lys Pro Gln Val Val Val Ala Pro Val Leu Met Ser Lys Leu 1 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 547:

(A) LENGTH: 367 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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35

30

Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Asn Pro Ser Gly Pro Val Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 35 40 45

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20

Tyr Gln Lys Lys Pro Val His Glu Lys Lys Glu Val Leu 65

INFORMATION FOR SEQ ID NO: 549: 3

(A) LENGTH: 47 amino acids SEQUENCE CHARACTERISTICS: Ξ 9

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 549: (D) TOPOLOGY: linear

2

. Met Leu Lys Gln Val Met Phe Val Phe Ser Gly Met Gly Pro Arg Ser l $_{\rm 1}$

His Cys Txp Gly Leu Pro Leu Ala Cys Gly Thr Phe Val Gln Gly His 20

2

Gln Ala Asp Ser Ser His Leu Leu Pro Leu Lys His Gln Gly Ala 35 40

(2) INFORMATION FOR SEQ ID NO: 550:

25

(A) LENGTH: 168 amino acids (i) SEQUENCE CHARACTERISTICS:

8

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 550:

Met Leu Leu Ser Leu Ala Ala Phe Ser Val Ile Ser Val Val Ser Tyr 35

Leu lle Leu Ala Leu Leu Ser Val Thr lle Ser Phe Arg lle Tyr Lys $$20\$ Gln Lys Ser Glu Glu Gly His Pro Phe Lys Ser Val Ile Gln Ala Val 6

Glu Ala Phe His Asn 60 Leu Asp Val Asp Ile Thr Leu Ser Ser 55 Ala Tyr 45

Leu 11e

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Asn Arg Ala

Ile Arg Leu Phe Leu Val Glu Asp Leu Val Asp Ser Leu Lys Leu Ala 95 Met Asn Ala Ala Met Val His Ile. S

Val Phe Met Trp Leu Met Thr Tyr Val Gly Ala Val Phe Asn Gly Ile 100 55

Tyr Glu Lys Tyr Lys Thr Gln Ile Asp His Tyr Val Gly Ile Ala Arg 130 Thr Leu Leu lle Leu Ala Glu Leu Leu Ile Phe Ser Val Pro Ile Val 115 8

Asp Gln Thr Lys Ser Ile Val Glu Lys Ile Gln Ala Lys Leu Pro Gly 145

s Lys Ala Glu Xaa 165 Ile Ala Lys Lys

S

INFORMATION FOR SEQ ID NO: 551: 2 9

(A) LENGTH: 124 amino acids (i) SEQUENCE CHARACTERISTICS: TYPE: amino acid

SEQUENCE DESCRIPTION: SEQ ID NO: 551: <u>e</u> Ŧ

2

Val Pro Phe His Leu Leu Val Val Leu Arg Ser Arg Ala Val Arg 5

Ala Arg Arg Arg Glu Pro Arg Ser Leu Pro Arg Pro Gly Asp Glu 25

2

Glu Leu Gln Leu Leu Cys Gly Ala Arg Ser Asp Phe Leu Glu Arg

23

Phe Pro Gly Ser Phe Gln Cys His Gln Cys Gly Phe Leu Pro His Pro 65 10 Cys Glu Glu Asp Trp Val Cys Leu Trp His His Ala Asp His Ala Ala 50 55 2

Gly Ser Ser Leu Cys His His Gln Leu Gln Asp Leu Gln Val Arg His 85 35

Ser Cys Thr Glu Val Arg Arg Pro Ser Ile Gln Ser Leu Pro 100 Pro

Gly Arg Arg His Tyr Ser Val Leu Arg Ser Phe Pro 115

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(2) INFORMATION FOR SEQ ID NO: 552:

45

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 177 amino acids TYPE: amino acid 3

SEQUENCE DESCRIPTION: SEQ ID NO: 552: æ

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Met Val His Leu Leu Val Leu Ser Gly Ala Trp Gly Met Gln Met Trp

Val Thr Phe Val Ser Gly Phe Leu Leu Phe Arg Ser Leu Pro Arg His 20 Thr Phe Gly Leu Val Gln Ser Lys Leu Phe. Pro Phe Tyr Phe His Ile 45 ŝ

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20 \$ 6 35 30 25 Ser Met Gly Cys Ala Phe Ile Asn Leu Cys Ile Leu Ala Ser Gln His 50 55 Ala Trp Ala Gln Leu Thr Phe Trp Glu Ala Ser Gln Leu Tyr Leu Leu 65 70 75 Arg Gln Leu Arg Glu Lys Asp Pro Lys Tyr Ser Ala Leu Arg Gln Asn $130\,$ Gly Leu Gly Gly Glu Val Pro Gly Ser His Gln Gly Pro Asp Pro Tyr 115 120 125 Arg Thr Thr Ala Ala Met Trp Ala Leu Gln Thr Val Glu Lys Glu Arg $105\,$ Phe Leu Ser Leu Thr Leu Ala Thr Val 85 Phe Phe Arg Tyr His Gly Leu Ser Ser Leu Cys Asn Leu Gly Cys Val 145 150 Leu Leu Gln Asn Cys Phe Glu Leu Leu Arg Thr Ser Thr Ser Gln Cys 35 40 Met Ala Phe Ile Leu Leu Phe Tyr Cys Leu Met Thr Phe Leu Ser Leu 1 5 (2) INFORMATION FOR SEQ ID NO: 553: ē Leu Ser Asn Gly Leu Cys Leu Ala Gly Leu Ala Leu Glu Ile Arg Ser 165 170 175 Thr Glu Gly Ile Pro Cys Ala Lys Ile Pro Glu Trp Val Thr His Leu 50 55 60 ž (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 553: (A) LENGTH: 72 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Asn Ala Arg Trp Leu Glu Pro 90 95

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Glu Gln Asn Ser Ala Thr Val Glu Pro Ser Ser His Glu Ile Leu His 20 25 10

Thr Trp Gln Thr Leu Lys Asn Ser 65 70

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(2) INFORMATION FOR SEQ ID NO: 554:

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(a) LENGTH: 45 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 554:

Val Glu Lys Ala His Asp Ser His Ser Ala Asp Pro Val Cys Pro Gly
25 30 Val Leu Arg Ile Ile Cys Leu Trp Pro Cys Gly Thr Thr Leu Pro Leu 1 15

5 Leu Thr Ala His Leu Pro Val Leu Leu Tyr Val Gln Leu 35 40 45

(2) INFORMATION FOR SEQ ID NO: 555:

5

(i) SEQUENCE CHARACTERISTICS (D) TOPOLOGY: linear (A) LENGTH: 251 amino acids (B) TYPE: amino acid

20 Met Lys His Ala Asp Pro Arg Ile Gln Gly Tyr Pro Leu Met Gly Ser 1 5 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 555:

30 25 Gly Phe Met Ile Val Tyr Asn Phe Ser Leu Val Ala Leu Ser Leu Tyr 50 55 Ser Leu Gly Pro Arg Ile Met Ala Asn Arg Lys Pro Phe Gln Leu Arg 35 40 45 Pro Leu Leu Met Thr Ser Ile Leu Leu Thr Tyr Val Tyr Phe Val Leu 20 25

35 Arg Cys Asp Pro Gln Asp Cys Thr Leu Gly Gln Cys Pro Ser Val Pro 85 90 95 Ile Val Tyr Glu Phe Leu Met Ser Gly Trp Leu Ser Thr Tyr Thr Trp 65 70 75 80

Ser Pro Xaa Thr Pro Val Thr Lys Ala Tyr Val Val Arg Thr Glu Gln \$100\$

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Trp Phe Leu Thr His Phe Pro Arg Ala Ala Pro Gly Met Trp Pro His 130 $$135\$ Gly Thr Gly Pro Pro Leu Pro Thr Ala Ala Leu Gln Gly Pro Arg Leu 115 120 125

3

Cys Cys Leu Pro Leu Gln Ser Trp Gly Leu Lys Gly Leu Tyr Ser Tyr
145 150 150

50

Phe Pro Leu Pro Ala Leu Lys Leu Gly Arg Gly Ala Leu Arg Ala Gly
175 Thr Lys Gly Leu Val Ala Phe Phe Leu Thr Gln Lys Arg Ser Ala 180 185

Ile Met Ser Leu Trp Thr Gln Ser His Ser Ser Thr Pro His Thr Glu
195 200 205

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Asp 240 Ala Val Ala Ser Gly Pro Lys Val Arg Val Gly Gly Gly Leu Gly Ile 210 120 Gin Pro Val Giu Ala Ala Tyr Ser Thr Cys Val Leu Ile Lys Ser 235

Arg Gly Asn His Glu Lys Lys Lys Lys Lys Lys 250 250

(2) INFORMATION FOR SEQ ID NO: 556:

2

SEQUENCE CHARACTERISTICS 3

2

(A) LENGTH: 19 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 556: (D) TOPOLOGY: linear

Leu Ala Gly Leu Cys Gly Gln Leu Ser Ser Pro Ala Leu Cys Val 5 Ğ ನ

Asn Arg Leu

25

(2) INFORMATION FOR SEQ ID NO: 557;

(A) LENGTH: 217 amino acids SEQUENCE CHARACTERISTICS: 3

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 557:

Met ile Thr Glu Lys Trp Gly Leu Asn Met Glu Tyr Cys Arg Gly Gln

35

Val Val Ala 30 Ę Ser Lys Met Phe Ser 25 Ala Tyr Ile Xaa Ser Ser Gly 6

Ser Arg Leu Leu Glu Lys Tyr Pro Gln Ala Ile Tyr Thr Leu Cys Ser 35 Trp Leu Ala Lys Ser Val Pro Val Met Gly 55 Ser Cys Ala Leu Asn Met 50 5

Ser Val Ala Leu Gly Thr Ile Glu Glu Val Cys Ser Phe Phe His 70 70 75 S & S Pro Gln Leu Leu Glu Leu Asp Asn Val Ile Ser Val Leu 95 Phe Gln Asn Ser Lys Glu Arg Gly Lys Glu Leu Lys Glu Ile Cys His 110 55

Arg Ser

Ser Gin Trp Thr Gly Arg His Asp Ala Phe Glu Ile Leu Val Glu Leu 115

Leu Gln Ala Leu Val Leu Cys Leu Asp Gly Ile Asn Ser Asp Thr Asn 9

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135

130

Tyr Ile Ala Gly Arg Ala Phe Val Leu Cys Ser 150 150 160 Ile Arg Trp Asn Asn 145

Phe Asp Phe Ile Val Thr Ile Val Val Leu Lys Asn 165 Ala Val Ser Asp

Val Leu Ser Phe Thr Arg Ala Phe Gly Lys Asn Leu Gln Gly Gln Thr 180

9

Ser Asp Val Phe Phe Ala Ala Gly Ser Leu Thr Ala Val Leu His Ser 195

Leu Asn Glu Val Ile Gly Lys Tyr Xaa 210

13

(2) INFORMATION FOR SEQ ID NO: 558: 2

(i) SEQUENCE CHARACTERISTICS;

(A) LENGTH: 82 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 558:

Leu Leu Lys Val Leu Cys Ile Leu Pro Val Met Lys Val Glu Asn Glu

Arg Tyr Glu Asn Gly Arg Lys Arg Leu Lys Ala Tyr Leu Arg Asn Thr

8

Leu Thr Asp Gln Arg Ser Ser Asn Leu Ala Leu Leu Asn Ile Asn Phe 45 Leu Asp Leu Met Val Asp Thr Tyr Ile Lys Leu 55 60 Ile Lys His Asp 50 Asp

35

Tyr Thr Ser Lys Ser Glu Leu Pro Thr Asp Asn Ser Glu Thr Val Glu 65 6

Asn Thr

45

(2) INFORMATION FOR SEQ ID NO: 559

(A) LENGTH: 95 amino acids SEQUENCE CHARACTERISTICS:

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(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 559; ê

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Met Val Leu Ile Leu Leu Asn Leu Leu Leu Gly Gln Phe Ser Cys Met 1 15 15

Ser Pro Ala Ser His His Cys His Pro Leu Pro Thr Glu Met Pro Cys 8

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Asp Ala Leu Leu Pro Lys Pro Ser Ala Asn Ser 50 55 Ser Ser Asp Txp Gly Phe Asp Ser His Thr Val Tyr Pro Ser Cys Val
35
40
45 Ala Glu Gly Pro Ala Ser Leu Arg Cys Asn Lys Tyr Val Ser 90 Gly His Pro Cys Val Gly Phe Ala Ala Val Leu Val Ala Pro Leu Thr \$40\$Cys His Cys Gin Gly Leu Tyr Asn Gin Gin Gin Gin Asn Leu His Ala 65 70 75 80 Tyr Tyr Pro Glu Ser Gly Trp Asp Arg Leu Arg Glu Leu Phe Gly Lys 50 55 60 Met Glu Val Pro Pro Pro Ala Pro Arg Ser Phe Leu Cys Arg Ala Leu 1 15 (2) INFORMATION FOR SEQ ID NO: 561: Val Ala Val Ser Ser Xaa 50 val Phe Thr Val Ile Gly Asp Ala Pro Gly Ala Val Leu Ser Cys Ala $20 \hspace{1cm} 25$ Met Ile Pro Ala Tyr Ser Lys Asn Arg Ala Tyr Ala Ile Phe Phe Ile 1 15 (2) INFORMATION FOR SEQ ID NO: 560: Glu Val Asp Thr Val Asn Thr Ser Leu Asn Val Tyr Arg Asn Lys Asp Ala Leu 65 70 75 Leu Glu Glu Arg Gln Lys Arg Leu Pro Tyr Val Pro Glu Pro 35 40 45 Ě E E SEQUENCE CHARACTERISTICS SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 561: SEQUENCE DESCRIPTION: SEQ ID NO: 560: (A) LENGTH: 108 amino (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 54 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear LENGTH: 108 amino acids Phe Pro Asn Gly Ser 60

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Ser His Phe Val Ile Ala Gly Ala Val Thr Gly Ser Leu Phe Arg Ile 85 95

Asn Val Gly Leu Arg Gly Trp Trp Leu Val Ala Xaa 100 105

5 (2) INFORMATION FOR SEQ ID NO: 562:

(1) SEQUENCE CHARACTERISTICS

20 15 Glu Gln Val Phe Leu Val Ile Leu Ala His Val Val Arg Arg Cys Ala 20 25 30 Met Asn Trp Gly Leu Ser Ile Trp Leu His Tyr Tyr Glu Lys Lys Lys 1 5 10 Ě (A) LENGTH: 50 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 562:

25 Ser Asp Gly Ile Leu Gln Phe Glu Ser Ser Leu Leu Lye Met Arg Arg 35

30 Ala Pro 50

(2) INFORMATION FOR SEQ ID NO: 563: (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 253 amino acids

3

6 Met Val Lys Val Cys Asn Asp Ser Asp Arg Trp Ser Leu Ile Ser Leu . 1 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 563:

Arg Arg Gin Phe Glu Phe Ser Val Asp Ser Phe Gin Ile Lys Leu Asp 35 $40\,$ Asn Asn Ser Gly Lys Asn Val Glu Leu Lys Phe Val Asp Ser Leu $20 \ 25 \ 30$

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Phe His Pro Thr Ile Ile Gly Glu Ser Val Tyr Gly Asp Phe Gln Glu 65 70 75 80 Leu Leu leu Phe Tyr Glu Cys Ser Glu Asn Pro Met Thr Glu Thr 50 $$5^\circ$$

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Glu Ile Arg Gly Gly Gly Leu Leu Lys Tyr Cys 100 105 Ala Phe Asp His Leu Cys Asn Lys Ile Ile Ala Thr Arg Asn Pro Glu 95 Asn Leu Leu Val Arg 110

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Gly Phe Arg Pro Ala Ser Asp Glu Ile Lys Thr Leu Gln Arg Tyr Met

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Cys Ser Arg Phe Phe Ile Asp Phe Ser Asp Ile Gly Glu Gln Gln Arg 130 120 115

Lys Tyr Glu Tyr Leu Met Thr Leu His Gly Val Val Asn Glu Ser Thr 175 Lys Leu Glu Ser Tyr Leu Gln Asn His Phe Val Gly Leu Glu Asp Arg 145

Met Gly His Glu Arg Arg Gln Thr Leu Asn Leu Ile Thr 180 Val Cys Leu

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Met Leu Ala Ile Arg Val Leu Ala Asp Gin Asn Val Ile Pro Asn Val 195 Ala Asn Val Thr Cys Tyr Tyr Gln Pro Ala Pro Tyr Val Ala Asp Ala 210 2

Asn Phe Ser Asn Tyr Tyr Ile Ala Gln Val Gln Pro Val Phe Thr Cys 225

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Gin Gin Gin Thr Tyr Ser Thr Trp Leu Pro Cys Asn Xaa 250

22

(2) INFORMATION FOR SEQ ID NO: 564: 2

(A) LENGTH: 18 amino acids SEQUENCE CHARACTERISTICS:

(B) TYPE: emino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 564:

33

Met Ser Phe Leu Met Trp Leu Met Ser Leu Ala Ile Thr Ser Gln Pro

Pro Met \$

INPORMATION FOR SEQ ID NO: 565: 3 45

(A) LENGTH: 80 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 565 (D) TOPOLOGY: 1inear

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Met Ala Pro Lys Gly Lys Val Gly Thr Arg Gly Lys Lys Gln Ile Phe 1 1 10 55 Ala Asn Ala Ile Tyr Cys Leu Val Thr Leu Val Phe Phr Ser Ser Ser 35 8

Glu Glu Asn Axg Glu Thr Leu Lys Phe Tyr Leu Axg Ile Ile Leu Gly 20

Ala Ser Phe Trp Ala Trp Leu Ala Leu Gly Phe Ser Leu Ala Val Tyr 50

Gly Ala Ser Tyr His Ser Met Ser Ser Met Ala Arg Ala Ala Phe Phe 65

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INFORMATION FOR SEQ ID NO: 566: (2)

(A) LENGTH: 73 amino acids SEQUENCE CHARACTERISTICS:

15

TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 566: (D) TOPOLOGY: linear

His Leu Lys Asp Val Ile Leu Leu Thr Ala Ile Val Gln Val Leu Ser

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Cys Phe Ser Leu Tyr Val Trp Ser Phe Trp Leu Leu Ala Pro Gly Arg 20 35

23

Ala Leu Tyr Leu Leu Trp Val Asn Val Leu Gly Pro Trp Phe Thr Ala

Asp Ser Gly Thr Pro Ala Pro Glu His Asn Glu Lys Ary Gln Ary Ary S0 60 9

Gln Glu Arg Arg Gln Met Lys Arg Leu 65

35

(2) INFORMATION FOR SEQ ID NO: 567:

SEQUENCE CHARACTERISTICS: 3

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(A) LENGTH: 263 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 567:

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Met Asp Cys Pro Ala Leu Pro Pro Gly Trp Lys Lys Glu Glu Val Ile 1 5

Gly Leu Ser Ala Gly Lya Ser Asp Val Tyr Tyr Phe Ser 20 30 Arg Lys Ser

S

Ser Gly Lys Lys Phe Arg Ser Lys Pro Gln Leu Ala Arg Tyr Leu 35 40 45

Lys Met Gly Asn Thr Val Asp Leu Ser Ser Phe Asp Phe Arg Thr Gly 22 Met Pro Ser Lys Leu Gln Lys Asn Lys Gln Arg Leu Arg Asn Asp Pro 65

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Leu Arg Gln Ser Gln Gly Lys Ser Pro Leu Leu Trp Lys Arg Thr Leu $20 \ 25 \ 30$ Oln Gly Val Gly Pro Gly Ser Asn Asp Glu Thr Leu Leu Ser Ala Val $165\,$ Met Met Axg Pro Phe Tyx Leu Leu Leu Pro Val Leu Cys 1 $^{\circ}$. 5 Ala Ser Ala Leu His Thr Ser Ser Ala Pro Ile Thr Gly Gln Val Ser 180 Asp Val Thr Glu Gln Ile Ile Lys Thr Met Glu Leu Pro Lys Gly Leu 145 150 150 Pro Arg Gln Leu Phe Trp Glu Lys Arg Leu Gln Gly Leu Ser Ala Ser 130 135 Leu Phe Gly Leu Thr His Leu 35 Glu Arg Val Gln Gln Val Arg Lys Lys Leu Glu Glu Ala Leu Met Ala 225 230 230 Ę Ala Ala Val Glu Lys Asn Pro Ala Val Trp Leu Asn Thr Ser Gln Pro 195 200 205 ďg (2) INFORMATION FOR SEQ ID NO: 568: Asp Ile Leu Cys Lys Ala Phe Ile Val Thr Asp Glu Asp Ile Arg Lys Oln Glu 210 \$215\$Ser Gly Asp Glu Ala Xaa 260 ž (i) SEQUENCE CHARACTERISTICS: Ser Arg Ala Ala Asp 245 SEQUENCE DESCRIPTION: SEQ ID NO: 568: (D) TOPOLOGY: linear B (A) LENGTH: 70 amino acids TYPE: amino acid Asn Pro Ser Ala Lys Leu Leu Ser 40 45 Thr Glu Glu Met Asp Ile Glu Met 250 255 e Thr Gln Ala 15

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His Pro Ser Asn Lys Val Lys Ser Asp Pro Gln Arg Met Asn Glu Gln 115 120 125 βıΑ Leu Asn Gln Asn Lys Gly Lys Pro Asp Leu Asn Thr Thr Leu Pro Ile 85 90 95 Gln Thr Ala Ser Ile Phe Lys Gln Pro Val Thr Lys Val Thr Asn $100\,$

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Gln Met Lys Thr Ser Gly Asn Arg Lys Ser Glu Tyr Ser Lys Tyr Ala 50 55 Asn Trp Lys Lys His 1

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Leu Leu

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Tyr Asp Thr

Pro Pro Pro Asp Cys His Cys His Ser Phe Arg Ala Glu 20 25 30

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 569:

(A) LENGTH: 34 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Pro Val Thr Ser Lys Arg Thr Leu 1

Phe Phe Pro Asp Pro Cys Ser

INFORMATION FOR SEQ ID NO: 570:

25

Ξ ž SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 570: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 104 amino acids

Met Asn Ser Arg Gly Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu 1 15

8

Trp Thr Leu Thr Asn Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu $35 \ 40 \ 45$ Leu His Ile Val Leu Leu Ser Ile Pro Phe Phe Ser Val Pro Val Ala 20 25 30

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His Ala Val Lys Gly Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ser 50 60

6

Ile Phe Thr Glu Val Phe His Asn Phe Ser Asn Asn Ser Ile Phe Ser 95 Ala Pro Asn Ser Leu Gly Thr Thr Gly Leu Trp Ser Thr Val Tyr
70 75 80

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Gly Lys Phe Leu Tyr Glu Val Xaa 100

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2 INFORMATION FOR SEQ ID NO: 571:

55

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 132 amino acids (B) TYPE: amino acid

ξ SEQUENCE DESCRIPTION: SEQ ID NO: 571: (D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 569:

(i) SEQUENCE CHARACTERISTICS:

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Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu Leu His Ile Val Leu $1 \ \ \, 10 \ \ \,$

Leu Ser Ile Pro Phe Ber Val Pro Val Ala Trp Thr Leu Thr Asn 20

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Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu His Ala Val Lys Gly 35

Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ala Arg Leu Leu Thr His 50 60

2

Trp Glu Gln Leu Asp Tyr Gly Val Gln Phe Thr Ser Ser Arg Lys Phe 65 Phe Thr Ile Ser Pro Ile Ile Leu Tyr Phe Leu Ala Ser Phe Tyr Thr 95

15

Pro Thr His Phe Ile Leu Asn Thr Ala Ser Leu Leu Ser 100 Lys Tyr Asp 20

Val Leu Ile Pro Lys Met Pro Gln Leu His Gly Val Arg Ile Phe Gly 120

ile Asn Lys Tyr 130

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(2) INFORMATION FOR SEQ ID NO: 572

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 32 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

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Leu Leu Ser Met Asn Lys Trp Ile Cys Glu Met His Cys Tyr Leu Val 1 5

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 572.

Val Cys Ser Pro Ser Ala Leu Arg Arg Val Arg His Thr Leu Ser Arg 20 20

5

(2) INFORMATION FOR SEQ ID NO: 573: တ

(A) LENGTH: 28 amino acids

23

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 573: (D) TOPOLOGY: linear

Met Pro Val Leu Ser Leu Leu Cys Thr Leu Ile Val Ser Phe Gln Ser 1 10 15

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Ala Asp Ser Cys Glu Val Phe Leu Asn Cys Ser Leu 25

(2) INFORMATION FOR SEQ ID NO: 574:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 40 amino acids (B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 574:

Met Lys Val Ser Thr Met Leu Trp Phe Leu Cys Trp Glu Gln Ser His $_{\rm 1}$

15

Phe Leu Arg Glu Trp Glu Asp Leu Ser Thr Phe Leu Ile Leu Ile Gln

Met Glu Cys Gln Tyr Gly Asn Ser 35

20

(2) INFORMATION FOR SEQ ID NO: 575: 22

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 30 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQUENCE 575.

8

Pro Ala Ser Met Gly Leu Pro Leu Met Ala Leu Met Trp Ser Thr 1 5

Ala Gly Val Asn Phe Ile Leu Ala Leu Pro Leu Leu Xaa 20

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(2) INFORMATION FOR SEQ ID NO: 576:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 29 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 576:

Met Lys Arg Gly Cys Leu Gly Leu Leu Phe Phe Ser Cys Cys Ser Ser 1 5

20

Ala Pro Thr Met Leu Leu Cys Asp Tyr Leu Asn Trp Phe 20

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(2) INFORMATION FOR SEQ ID NO: 577:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 577: (B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Lys Leu Leu Gly Ile Ala Leu Leu Ala Tyr Val Ala Ser Val 1 15

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Lys Ile Glu Ser Lys Ile Glu Glu Met Val Glu Pro Leu Arg Glu Lys \$45\$Trp Gly Asn Phe Val Asn Met Arg Ser Ile Gln Glu Asn Gly Glu Leu $20 \hspace{1cm} 25$

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Ile Arg Asp Leu Glu Lys Ser Phe Thr Gln Lys Tyr Pro Pro Val Lys $50 \ \ 55$

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Phe Leu Ser Glu Lys Asp Arg Lys Arg Ile Leu Xaa Asn 65 70 75 n Arg Arg Arg 80

XAA Val Arg Gly Leu Pro Ser Xaa Leu Thr Asn Ser 85 90

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(2) INFORMATION FOR SEQ ID NO: 578:

25

E SEQUENCE CHARACTERISTICS (A) LENGTH: 42 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

30

SEQUENCE DESCRIPTION: SEQ ID NO: 578:

Phe Ser Leu Val Leu Leu Ile Lys Ile Ile Ser Phe Glu Arg 5 10 15

Met Lys

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ž Leu Ile Phe Leu Phe Pro Leu Ser Phe Leu Pro Asn Ile Trp Arg 20 25 30

6 ρıς Va. Met Val Asn Leu Asn Ile Leu Phe 35

(2) INFORMATION FOR SEQ ID NO: 579:

45

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 70 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 579:

S

Leu Ala Gin Glu Cys Pro Pro His Ile Pro Ser Ser Phe Phe Leu Val 1 10 , 5

Lyg Leu Leu Phe Ile Pro Trp Leu Ala Ser Leu Leu Pro Pro Leu Ser $20 \ 30$

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Thr Phe Thr Ser Asp Phe Tyr Phe Met Glu Phe Gly Ile Glu Val Lys $_{1}^{1}$ $_{2}^{1}$

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Leu Gln Gln Cys Arg Gln His Gln Val Leu Gln Glu Lys Asn Thr Lys 50 60

Lys Phe Asn Lys Lys Lys 65 70

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INFORMATION FOR SEQ ID NO: 580:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 110 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 580:

5

Met Leu Arg Leu Leu Leu Val Ala Phe Ala Leu Val Val Val Leu 1 10 15 Phe His Val Leu Leu Ala Pro Ile Thr Ala Leu Phe His Thr His Phe $20 \ 25 \ 30$

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Tyr Asp Arg Leu Gin Asp Ala Gly Ser Arg Trp Pro Glu Leu Tyr Leu 35

25

Tyr Ser Arg Ala Asp Glu Val Val Leu Ala Arg Asp Ile Glu Arg Met 50 55 60

Val Glu Ala Arg Leu Ala Arg Arg Val Leu Ala Arg Ser Val Asp Phe 65 70 75

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Val Ser Ser Ala His Val Ser His Leu Arg Asp Tyr Pro Thr Tyr Tyr Tyr 95

Thr Ser Leu Cys Val Asp Phe Met Arg Asn Cys Val Arg Cys
100 105 110

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6 (2) INFORMATION FOR SEQ ID NO: 581:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 30 amino acids

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(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 581:

Met Phe Lys Leu Glu Glu Cys Gly Lys Thr Thr Phe Leu Leu Ser Met
1 10 15

Ala Leu Tyr Phe Trp Trp Ile Val Gln Thr Thr Lys Gly Cys
25 30

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2 INFORMATION FOR SEQ ID NO: 582:

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(1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 71 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 582: (D) TOPOLOGY: linear

Met Glu Ser Asp Ala Leu Leu Leu Thr Ile Phe Trp Ile Ile Ala Arg

S

Ser Ser Val Arg Ser Val Gly Lys Ser Ser Gln Arg Ser Phe Thr Thr 20 Ile Thr Gln Leu Arg Ser Thr His Thr Gly Pro Ser Arg Arg Ser Tyr

2

Leu ile Trp Trp Asn Gly Gly Pro Lys Arg Thr ile Ser Tyr Val Ser 50 60

Arg Arg Phe Arg Ser Phe Arg 65 70

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(2) INFORMATION FOR SEQ ID NO: 583:

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(i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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val Gly Leu Phe Gln Pro Lys Thr Phe Gln Val Pro Val Thr Asp Leu

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 583:

Tyr ile Phe ile Lys ile Tyr Ser Glu ile Gly Pro ile Met His Val

Pro Gly Tyr Ser Gln Ser Pro Ser Thr Pro Pro Trp Thr Leu Cya 33

(2) INFORMATION FOR SEQ ID NO: 584; 6

SEQUENCE CHARACTERISTICS:

TYPE: amino acid

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SEQUENCE DESCRIPTION: SEQ ID NO: 584:

Met Trp Phe Gly Ser Asp Arg Ser Asp Leu Arg Ile Gly Thr Ala Phe 1 5 S

Leu Phe Asp Leu Val Cys Asp Leu Cys Ile His Ala Trp Lys Pro Pro

Gly Leu Val Arg Phe Ser Phe

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(2) INFORMATION FOR SEQ ID NO: 585:

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(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 19 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 585:

Leu Asn Thr Ala Ser Leu Asn Leu Pro Trp Lys Val Gln Leu Phe 10 15

Ala His Ala 9

(2) INFORMATION FOR SEQ ID NO: 586: 12

(A) LENGTH: 23 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 586:

Met Ser Ala Cys Leu Leu Leu Phe Leu Ala Phe Ser Trp Lys Arg Lys

Gly Leu Trp Ser Gly Pro Gly

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(2) INFORMATION FOR SEQ ID NO: 587:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 69 amino acids (B) TYPE: amino acid

33

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 587; (D) TOPOLOGY: linear

Leu Pro Pro Phe Ser Leu Val Tyr Thr His Phe Leu Val Ala Ser 10 Met

4

Leu Leu Pro Val Ile Leu Ala Val Phe Pro Asp Ser Ala Gln Ile Val

Pro Leu Lys Pro Ile Pro Arg Pro Gln Pro Glu Val Ile Phe Pro 35 40 5

Ser Ser Glu Leu Leu Glu Gln Leu Leu Ser Val Gln Phe Val Trp Gln 50 60

Ala His Thr Val Ala

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(2) INFORMATION FOR SEQ ID NO: 588:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 77 amino acids (B) TYPE: amino acid

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INFORMATION FOR SEQ ID NO: 590:

(1) SEQUENCE CHARACTERISTICS:

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ᅜ 5 25 20 55 **\$** 6 $\frac{3}{5}$ 30 S 8 ጀ Net Pro Glu Thr Arg Leu Gly His Arg Gln Gln Phe Ala Val Phe His 1 10 15 Met Leu Thr Phe Leu Phe Ser Ala Cys Ala Thr Cys Leu Gly Lys Leu 1 15 Met Asp Pro Phe His Tyr Asp Tyr Gln Thr Leu Arg Ile Gly Gly Leu 10 15 Gly Pro Pro Leu Leu Ser 35 Ala Ser Pro Leu Ala Pro Val Gly Pro Gln Gln Arg Gly Xaa Pro Pro 20 25 30 (2) INFORMATION FOR SEQ ID NO: 591: Glu Glu Ala Gln Val Glu Asn Leu Ile Thr Ala Asn Ala Thr Glu Pro 50 55 60 Arg Arg Cys Lys Cys Ser Phe Asn Gln Lys Pro Arg Ala Pro Gly Asp 45Val Phe Ala Val Val Leu Phe Ser Val Gly Ile Leu Leu Ile Leu
20 25 30 (2) INFORMATION FOR SEQ ID NO: 592: Xaa Pro Val Pro Pro Cys Gly Ě (1) SEQUENCE CHARACTERISTICS (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 591: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 592: (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 590: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 24 amino acids (A) LENGTH: 38 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 69 amino acids

Gln Leu Leu Ala,Gln Thr Lys Lys Val Val Arg Pro Thr Arg Lys Lys 130 135 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 589: (A) LENGTH: 155 amino e (B) TYPE: amino acid (D) TOPOLOGY: linear

Val Gly Phe Leu Glu Leu Leu Ala Gly Leu Leu Leu Val Met Gly Pro 65 70 75 Leu Lys Val Phe Gly Tyr Gln Pro Asp Pro Leu Asn Tyr Gln Ile Ala 50 55 Ser Glu Arg Met Asn Ala Leu 35

Phe Val Gln Phe Ala Glu Val Phe Pro 40

Met Ala Leu Leu Ser Val Leu Arg Val Leu Leu Gly Gly Phe Phe
1 10 15

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(2) INFORMATION FOR SEQ ID NO: 589:

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SEQUENCE CHARACTERISTICS

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Asn Val Gly Gln Leu Leu Ala Gln Thr Lys Lys Val Val Arg Pro Thr $50\,$

Arg Lys Lys Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys 65

Ser Thr Cys Ile Pro Ala Ile Val Cys Leu Gly Phe Leu Leu Leu Leu 35

Leu Met Met Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu 20 30

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Met Gly Pro Pro Met Leu Gln Glu Ile Ser Asn Leu Phe Leu Ile Leu 1 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 588:

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Pro Met Leu Gln Glu Ile Ser Asn Leu Phe Leu Ile Leu Leu Met Met . 95

Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu Ser Thr Cys 100 105

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Ile Pro Ala Ile Val Cys Leu Gly Phe Leu Leu Leu Leu Asn Val Gly
115 120 125

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Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys Xaa 145 150 150

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Gln Lys Ala Glu Asn 65

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WO 98/39448 PCT/US98/04493		(2) INFORMATION FOR SEQ ID NO: 593:	(i) SEQUENCE CHARACTERISTICS: (A) LENTH: 308 amino acids (B) TVPE: amino acids	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 593:	Asn Leu Arg Val Arg Leu Gly Asp Val Ile Ser Ile Gln Pro Cys Pro 1 5 10	Asp Val Lys Tyr Gly Lys Arg Ile His Val Leu Pro Ile Asp Asp Thr 20	Val Glu Gly Ile Thr Gly Asn Leu Phe Glu Val Tyr Leu Lys Pro Tyr 35 40	Phe Leu Glu Ala Tyr Arg Pro Ile Arg Lys Gly Asp Ile Phe Leu Val 50 55	Arg Gly Gly Met Arg Ala Val Glu Phe Lys Val Val Glu Thr Asp Pro 65 80	Ser Pro Tyr Cys Ile Val Ala Pro Asp Thr Val Ile His Cys Glu Gly 85	Glu Pro 11e Lys Arg Glu Asp Glu Glu Ger Leu Asn Glu Val Gly 100	Tyr Asp Asp ile Gly Gly Cys Arg Lys Gln Leu Ala Gln Ile Lys Glu 115 . 120	Met Val Glu Leu Pro Leu Arg His Pro Ala Leu Phe Lys Ala Ile Gly 130	Val Lys Pro Pro Arg Gly Ile Leu Tyr Gly Pro Pro Gly Thr Gly 145 160	Lys Thr Leu Ile Ala Arg Ala Val Ala Asn Glu Thr Gly Ala Phe 175	Phe Leu Ile Asn Gly Pro Glu Ile Met Ser Lys Leu Ala Gly Glu Ser 180	Glu Ser Asn Leu Arg Lys Ala Phe Glu Glu Ala Glu Lys Asn Ala Pro 195	Ala Ile Ile Phe Ile Asp Glu Leu Asp Ala Ile Ala Pro Lys Arg Glu 210	Lys Thr His Gly Glu Val Glu Arg Arg Ile Val Ser Gln Leu Leu Thr 225 240	Leu Met Asp Oly Leu Lys Gln Arg Ala His Val Ile Val Met Ala 245	Thir Asn Arg Pro Asn Ser Ile Asp Pro Ala Leu Arg Arg Phe Gly Arg 260	Phe Asp Arg Glu Val Asp Ile Gly Ile Pro Asp Ala Thr Gly Arg Leu

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 601: (B) TYPE: amino acid
(D) TOPOLOGY: linear

S Met Trp Thr Arg Ser Ser Arg Cys Leu Leu Ceu Cys Ile Pro Gly Xaa 1 15

5 Ser Arg Arg Arg Ala Gly Ser Gly Met Lys Pro Arg Ser Trp Ser 20 25 Ala Trp Arg Pro Ser Gly Gly Thr Gly Thr Ser Ser Ser Gln Ser Ser 35 40 45

2 The Gln Ser Arg The Leu Ser Ala The Ala Ser Pro Ala 50

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 8 amino acids

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(2) INFORMATION FOR SEQ ID NO: 598:

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SEQUENCE DESCRIPTION: SEQ ID NO: 597:

(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 1 amino acids

(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 602:

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(1) SEQUENCE CHARACTERISTICS: (A) LEWSTH: 29 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Arg Glu Thr Ser Ile Arg Val Leu Leu Met Leu Pro Ala Leu Glu 1 15 X SEQUENCE DESCRIPTION: SEQ ID NO: 602:

23

3 Ser Thr Ser Gly Leu Ser Ala Phe Met Gly Leu Gly Thr 20 25

ઝ (2) INFORMATION FOR SEQ ID NO: 603:

Œ Ê SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 603: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 69 amino acids

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Met Pro Pro Lys Gln Glu Leu Gly Ser Gly Val Gly Glu Leu Ala Lys

45

Asm Ser Lys Arg Gln His Trp Asm His Arg Trp Lys Lys Tyr Leu Lys 20 25 30

8 Leu Glu His Cys Ala Thr Met Ala Trp Asp Cys Leu Met Arg Leu Glu 50 \$5Leu Ile Arg Trp Glu Asp Gly Leu Leu Leu Glu Gly Leu Leu Leu Val
35 40 45

S Leu Leu Lys Arg Leu 65

INFORMATION FOR SEQ ID NO: 601:

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Met Xaa Gly Leu Ser Leu Ile Leu Thr Val Thr Leu Leu Ala Val Ser 1 10

Asp Ser Ala Ala Thr Cys Ile Val Ala Lys Gly
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SEQUENCE DESCRIPTION: SEQ ID NO: 600:

(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 27 amino acids 6

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 600:

35

Val His Thr Pro Ser Arg Leu Pro Ala 20

Met Phe Leu Val Trp Phe Phe Trp Gly Leu Ile Ser Ala Leu Ser Asn 1 5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 599:

(D) TOPOLOGY: linear (A) LENGTH: 25 amino acids

(B) TYPE: amino acid

30

25

(2) INFORMATION FOR SEQ ID NO: 599:

(i) SEQUENCE CHARACTERISTICS:

20

Met Cys Ile Met Ser Ala Leu Val

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SEQUENCE DESCRIPTION: SEQ ID NO: 598:

(B) TYPE: amino acid
(D) TOPOLOGY: linear

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 61 amino acids

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(2) INFORMATION FOR SEQ ID NO: 604:

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Lys lle Val Tyr lle Leu Gly Asn Pro Leu Lys Phe Asn Ser Arg Val 1 5 10 15

Ile His His Leu Val Leu Leu Gln 20

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 604:

(B) TYPE: amino acid (D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids

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23

Met Asn Leu His Gln Arg Arg Leu Leu Leu Ile Gly His Leu Met Thr

23

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 605:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 35 amino acids

(B) TYPE: amino acid

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(D) TOPOLOGY: linear

(2) INFORMATION FOR SEQ ID NO: 605:

15

Leu Val Lys Ala Ser Lys Ser Phe Ser Phe Thr Glu Ile Thr Ser Ser

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Met Ala Pro Pro Gly Trp Gln Xaa Xaa Xaa Xaa Trp Leu Ala Cys 1 10 11 Pro Asp Arg Gly Glu Leu Ser Ser Arg Ser Pro Pro Cys Arg Leu Ala 20 Arg Trp Ala Glu Gly Asp Arg Glu Thr Arg Thr Cys Leu Leu Glu Leu 15. Ser Ala Gln Ser Trp Gly Gly Arg Phe Arg Arg Ser Ser Ala Val Ser 50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 608: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 609: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 232 amino acids (A) LENGTH: 6 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 609: (B) TYPE: amino acid (D) TOPOLOGY: linear Glu Leu Asp Tyr Ile Leu 8 33 各 45 ಬ 25

Leu Leu Gly Tyr Gly Leu Phe Gly His Cys Ile Val Leu Phe Ile Thr $_{\rm 1}$ $_{\rm 10}$

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(b) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 606:

6

(A) LENGTH: 130 amino acids

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 606:

35

Lys 35

Arg Lys

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Tyr Asn Ile His Leu His Ala Leu Phe Tyr Leu Phe Trp Leu Leu Val 20

Gly Gly Leu Ser Thr Leu Arg Met Val Ala Val Leu Val Ser Arg Thr

20

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Glu Gly Ile Leu Asp Thr Leu Glu Gly Pro Asn Ile Pro Pro Ile Gln 85 85

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His Met Leu Phe Leu Leu Tyr Leu His Phe Ala Tyr His Lys Val Xaa 65 75 80

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Val Gly Pro Thr Gln Arg Leu Leu Cys Gly Thr Leu Ala Ala Leu 50 60

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Ala Gly Ser Pro Ser Arg Leu His Phe Leu Pro Gln Pro Leu Leu Leu 65 70 75 Arg Ser Ser Gly Ile Pro Ala Ala Ala Thr Pro Trp Pro Gln Pro Ala 95

Gly Leu Pro Val Arg Pro Thr Pro Thr Arg Thr Gly Glu Glu Asp Arg 100 100

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Thr Leu Asp Ile Ser Ile Cys Thr Glu Val Leu Ala Gly Thr Glu Gln 115 120 128

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Pro Pro Pro Pro Arg Met Thr Ser Pro Ser Ser Ser Pro Val Phe Arg 130 140

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Leu Glu Thr Leu Asp Gly Gly Gln Glu Asp Gly Ser Glu Ala Asp Arg 145 150 150

Gly Lys Leu ₽ B Phe Gly Ser Gly Leu Pro Pro Met Glu Ser Gln Phe 165 170 175

20

Gly Lys Lys Asn Gln Leu Leu Val Ile

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SEQUENCE DESCRIPTION: SEQ ID NO: 612:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 9 amino acids

25

(2) INFORMATION FOR SEQ ID NO: 613:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 29 amino acids

15

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(2) INFORMATION FOR SEQ ID NO: 612:

(i) SEQUENCE CHARACTERISTICS:

Pro Thr Ser His Pro 20

Met Val Phe Glu Gly Phe Ser Ser Ala Phe Cys Leu Ser Ser Thr Ala 1 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 611:

8

Gln Gly Glu Asp Arg Lys Phe Ala Pro Ser Asp Lys Ser Gln Pro Pro 180 180

Thr Thr Glu Arg Glu Gln Val Pro Val Ser Arg Ile Gln Thr Asp Leu 195 200 205

25

Thr Glu Ile Gly Ser Ser Met Arg Ser Pro Gly Val Ser Pro Arg Ile 210 225

30

Trp Leu Asp 225

Phe Gln Ser Thr Xaa 230

35

(2) INFORMATION FOR SEQ ID NO: 610:

(1) SEQUENCE CHARACTERISTICS:

ઝ Met Val Trp Val Leu Trp Ser Ala Pro Ser Leu Ala Pro Pro Trp Val
1 15 30

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 613:

(D) TOPOLOGY: linear (B) TYPE: amino acid

Gly Pro Cys Trp Pro Ser Thr Gly Asn Cys Cys Leu Cys
20 25

6 (2) INFORMATION FOR SEQ ID NO: 614:

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SEQUENCE DESCRIPTION: SEQ ID NO: 610:

(D) TOPOLOGY: linear (A) LENGTH: 34 amino acids
(B) TYPE: amino acid

(i) SEQUENCE CHARACTERISTICS:

Σ SEQUENCE DESCRIPTION: SEQ ID NO: 614: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 30 amino acids

3

Met Ala Lys Arg Ser Pro Gly Gly Cys Gly Ser Gly Leu Ile Leu Leu 1 15

Cys Cys Gln Pro Cys Arg Pro Thr Ser Ser Ala Pro Met Arg

8

(2) INFORMATION FOR SEQ ID NO: 615:

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(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 113 amino acids (B) TYPE: amino acid

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 21 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 611:

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Arg His

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Met Val Leu Leu Leu Leu Ala Tyr Val Leu Leu Thr Tyr Ile Leu 1 5

Leu Leu Asn Met Leu Ile Ala Leu Met Xaa Arg Asp Arg Gln Gln Cys 20 30

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 615: (D) TOPOLOGY: linear

Ile Thr Ile Ala Ile Gln Met Ile Cys Leu Val Asn Xaa Glu Leu Tyr

Pro Thr Phe Val Arg Asn Xaa Gly Val Met Val Cys Ser Ser Leu Cys 20

Asp lle Gly Gly Ile Ile Thr Pro Phe Ile Val Phe Arg Leu Arg Glu 35 45 2

Ę ž Leu Gly Leu Ile Leu Phe Ala Val 55 60 Val Trp Gln Ala Leu Pro 50

15

Leu Leu Leu Pro Glu Thr Lys Gly Val Ala Leu 70 75 80 Ala Ala Gly Val Thr 65

Pro Glu Thr Met

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Lys Glu Asn Thr 11e Tyr Leu Lys Val Gln Thr Ser Glu Pro Ser Gly 100 Lys Asp Ala Glu Asn Leu Gly Arg Lys Ala Lys 85

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(2) INFORMATION FOR SEQ ID NO: 616: 39

(i) SEQUENCE CHARACTERISTICS:

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 616:

Thr Met Lys Asp Ala Glu Asn Leu Gly Arg Lys Ala Lys Pro Lys Glu 1 1 15

Asn Thr

(2) INFORMATION FOR SEQ ID NO: 617:

45

(A) LENGTH: 21 amino acids SEQUENCE CHARACTERISTICS: 3

(B) TYPE: amino acid

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SEQUENCE DESCRIPTION: SEQ ID NO: 617: (D) TOPOLOGY: linear (xį Pro Arg Val Arg Asn Ser Pro Glu Asp Leu Gly Leu Ser Leu Thr Gly 1 5 5 10 55

Asp Ser Cys Lys Leu 20

8

(2) INFORMATION FOR SEQ ID NO: 618:

(A) LENGTH: 52 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 618:

cin Ala Asp Asp Leu Gin Ala Thr Val Ala Ala Leu Cys Val Leu Arg

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Pro Gly Gly Gly Gly Fro Trp Ala Gly Ser Trp Leu Ser Pro Lys Thr $20\ 25\$ Ala Met Gly Gly Asp Leu Val Leu Gly Leu Gly Ala Leu Arg Arg Arg 35

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Lys Arg Leu Leu 50

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(2) INFORMATION FOR SEQ ID NO: 619:

25

(1) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 619:

30

Glu Gln Glu Lys Ser Leu Ala Gly Trp Ala Leu Val Leu Ala Xaa Xaa 1 10 15

Cys Ser Ala Val Asn Ala Thr Gly His Leu Ser Asp Thr Leu Trp Leu Gly lle Gly Leu Met Val Leu His Ala Glu Met Leu Trp Phe Gly Gly 30 26

35

lle Pro lle Thr Phe Leu Thr lle Gly Tyr Gly Asp Val Val Pro Gly 50 50

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The Met Trp Gly Lys Ile Val Cys Leu Cys Thr Gly Val Met Gly Val 65 5

Cys Cys Thr Ala Leu Leu Val Ala Val Val Ala Arg Lys Leu Glu Phe $85\ \,$ 90

Asn Lys Ala Glu Lys His Val His Asn Phe Met Met Asp Ile Gln Tyr 100 S

Thr Lys Glu Met Lys Glu Ser Ala Ala Arg Val Leu Gln Glu Ala Trp 115 Met Phe Tyr Lys His Thr Arg Arg Lys Glu Ser His Ala Ala Arg Xaa 130 130

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His Gin Arg Xea Leu Leu Ala Ala Ilo Asn Ala Pho Arg Gin Val Arg 145

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Leu Lys His Arg Lys Leu Arg Glu Gln Val Asn Ser Met Val Asp Ile 165 170 175

S Ser Lys Met His Met Ile Leu Tyr Asp Leu Gln Gln Asn Leu Ser Ser 180 185

Ser His Arg Ala Leu Glu Lys Gin Ile Asp Thr Leu Ala Gly Lys Leu 195 200 205

5

Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly Pro Arg Gln Leu 210 215

5 Pro Glu Pro Ser Gln Gln Ser Lys 230

2 INFORMATION FOR SEQ ID NO: 620:

20

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 36 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 620:

25

Tyr Gln Ala His His Val Ser Arg Asn Lys Arg Gly Gln Val Val Gly
1 15

30 Thr Arg Gly Gly Phe Arg Gly Cys Thr Val Trp Leu Thr Gly Leu Ser $20\ 25\$

Gly Ala Gly Lys 35

35

3 INFORMATION FOR SEQ ID NO: 621:

Ê SEQUENCE CHARACTERISTICS: (A) LENGTH: 57 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 621: (B) TYPE: amino acid
(D) TOPOLOGY: linear

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Asn Tup His Met Lys Lys His Asp Ala Asp Ser Phe Tyr Gln Phe Ser 25 30 Leu Gln Cys Glu Ile Cys Gly Phe Thr Cys Arg Gln Lys Ala Ser Leu 1 5 10

Cys Asn Ile Cys Gly Lys Lys Phe Glu Lys Lys Asp Ser Val Val Ala $35\,$. $40\,$

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His Lys Ala Lys Ser His Pro Glu Val 50 55

(2) INFORMATION FOR SEQ ID NO: 622:

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(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 19 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 622:

Ile Thr Ser Thr Asp Ile Leu Gly Thr Asn Pro Glu Ser Leu Thr Gln
1 15

Pro Ser Asp

5

(2) INFORMATION FOR SEQ ID NO: 623:

7

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 19 amino acids

(B) TYPE: amino acid

20

X SEQUENCE DESCRIPTION: SEQ ID NO: 623: (D) TOPOLOGY: linear

Asn Ser Thr Ser Gly Glu Cys Leu Leu Clu Ala Glu Gly Met Ser 1 1 5 .

Lys Ser Tyr

25

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(2) INFORMATION FOR SEQ ID NO: 624:

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 51 amino acids

35

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 624:

Cys Ser Gly Thr Glu Arg Val Ser Leu Met Ala Asp Gly Lys Ile Phe 1 5 10

8

Val Gly Ser Gly Gly Gly Thr Glu Gly Leu Val Met Asn Ser
20 25 30

Asp Ile Leu Gly Ala Thr Thr Glu Val Leu Ile Glu Asp Ser Asp Ser 35 40 45

25

Ş Ala Gly Pro 50

(2) INFORMATION FOR SEQ ID NO: 625:

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(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 60 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 625: (D) TOPOLOGY: linear

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Ile Gin Tyr Val Arg Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala 1 5 15 Lys Lys Tyr Val Cys Pro His Pro Ser Cys Gly Arg Leu Phe Arg Leu 15 His Pro Arg Tyr Leu Gln His His Ile Lys Tyr Gln His Leu Leu Lys $20\ 20$ Asp Gln Arg Asp Tyr Ile Cys Glu Tyr Cys Ala Arg Ala Phe Lys Ser 1 5 Arg Ser Ser Arg Ser Lys Thr Gly Ser Leu Gln Leu Ile Cys Lys Ser 1 5 10 Leu Gln Cya Glu Ile Cya Gly Phe Thr Cys Arg Gln Lys Ala Ser Leu Asn Trp His Met Lys Lys His Asp Ala Asp Ser Phe Tyr Gin Phe Ser 20 Ser His Asn Leu Ala Val His Arg Met Ile His Thr Gly Glu Lys 20 30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 626: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 627: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 628: Gln Lys Gln Leu Leu Arg His Ala Lys His His Thr 50 55 (A) LENGTH: 31 amino acids (A) LENGTH: 25 amino acids (i) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear Glu Pro Asn Thr Asp Gln Leu Asp Tyr 20 (2) INFORMATION FOR SEQ ID NO: 626: (2) INFORMATION FOR SEQ ID NO: 627: (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 628: 3 Ξ 15 2 22 35 3 တ္တ 8 45 55

Lys Tyr Val Cys Pro His Pro Ser Cys Gly Ary Leu Phe Ary Leu Gln 135 46 : Lys Ser 95 Arg Ala Phe Lys Ser Ser His Asn Leu Ala Val His Arg Met Ile His 175 Gin Tyr Val Arg Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala His $_{\rm 1}$ Pro Arg Tyr Leu Gln His His Ile Lys Tyr Gln His Leu Leu Lys Lys 20 25Asp Ile Leu Gly Thr Asn Pro Glu Ser Leu Thr Gln Pro Ser Asp Xea Asn Ser 65 70 70 Tyr Xaa Cys Ser Gly Thr Glu Arg Val Ser Leu Met Ala Asp Gly Lys 100 lie Phe Val Gly Ser Gly Ser Gly Gly Thr Gly Gly Leu Val Met 115 Ser Asp lie Leu Gly Ala Thr Thr Glu Val Leu lie Glu Asp Ser 130 Asp Ser Ala Gly Pro Xea Gln Arg Asp Tyr Ile Cys Glu Tyr Cys Ala 145 Cys Asn Ile Cys Gly Lys Lys Phe Glu Lys Lys Asp Ser Val Val Ala 35 His Lys Ala Lys Ser His Pro Glu Val Xaa Ile Thr Ser Thr Thr Ser Gly Glu Cys Leu Leu Leu Glu Ala Glu Gly Met Ser (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 629: Lys Gln Leu Leu Arg His Ala Lys His His Thr Asp 50 55 (A) LENGTH: 60 amino acids SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 629: (2) INFORMATION FOR SEQ ID NO: 630: Thr Gly Glu Lys His Tyr Xas 180 Э Asn 8 35 6 45 S 15 ឧ 23 25 S 2

(A) LENGTH: 27 amino acids (B) TYPE: amino acid

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(2) INFORMATION FOR SEQ ID NO: 631:

(1) SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 634:

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Asp Leu

Val His Arg Glu Glu Ala Ser Cys Tyr Cys Gln Ala Glu Pro Ser Gly 1 15

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(A) LENGTH: 110 amino acids (B) TYPE: amino acid

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Pro Phe Lys Asp Asp Pro Arg Asp Glu Thr Tyr Lys Pro His Leu Glu 1 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 630:

(D) TOPOLOGY: linear

Arg Glu Thr Pro Lys Pro Arg Arg Lys Ser Gly
20 25

25

Leu Tyr Leu His Asn His Ser Tyr Ala Ser Ile Gln Ser Asp Asp Leu 35 40

Trp Asp Ser Phe Asn Glu Val Thr Asn Gln Thr Leu Asp Val Lys Arg 50 55

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Gln Lys Lys Gly Lys Glu Leu Phe Ile Gln Gln Glu Arg Phe Phe Leu 95

Met Met Lys Thr Trp Thr Leu Gln Lys Gly Phe Pro Leu Val Thr Val 65 70 75

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Σ

SEQUENCE DESCRIPTION: SEQ ID NO: 631:

(D) TOPOLOGY: linear

Met Leu Lys Thr Tyr Leu Ser Glu Asp Val Phe Gln His Ala Val Val 20

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ၓ 25 20 45 6 35 50 S Lys Arg Trp Ala Gly Leu 20 Arg Pro Ala Leu Arg Gln Ala Gly Gly Gly Thr Arg Glu Pro Arg Gln
1 15 Ser Leu Phe Leu Leu Val Val Leu Tyr His Tyr Val Ala Val Asn Asn 20 25 30 Met Ile Thr Asp Val Gin Leu Ala Ile Phe Ala Asn Met Leu Gly Val 1 15 (2) INFORMATION FOR SEQ ID NO: 636: Ala Val Asn Phe Arg Pro Gln Arg Ser Gln Ser Met (2) INFORMATION FOR SEQ ID NO: 635: Pro Lys Lys Gln Glu 35 (1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 635: (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 634: Œ (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 636: (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 22 amino acids (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 37 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 12 amino acids

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Asn

Met Lys Pro Glu Ile Gln Pro Ser Asp Thr Arg Tyr Met 100 105 110

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Leu Glu Lys Val Ala Ser Val Gly Asn Ser Arg Pro Thr Gly Gln Gln 1 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 632:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 24 amino acids

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Leu Glu Ser Leu Gly Leu Leu Ala 20

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(2) INFORMATION FOR SEQ ID NO: 633:

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INFORMATION FOR SEQ ID NO: 632:

(1) SEQUENCE CHARACTERISTICS:

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(1) SEQUENCE CHARACTERISTICS:

(B) IENGTH: 18 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 633:

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INFORMATION FOR SEQ ID NO: 637: 3

(A) LENGTH: 342 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS

SEQUENCE DESCRIPTION: SEQ ID NO: 637: (D) TOPOLOGY: linear Œ

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Glu Glu Met Ala Asp Ser Val Lys Thr Phe Leu Gln Asp Leu Ala Arg 1 15

Gly lie Lys Asp Ser lie Trp Gly lie Cys Thr lie Ser Lys Leu Asp \$20\$

Ala Arg Ile Gin Gin Lys Arg Giu Giu Gin Arg Arg Arg Ala Ser 15

15

Val Leu Ala Gin Arg Arg Ala Gin Sex Ile Giu Arg Lys Gin Glu 50 60 Ser

2

118 Glu Pro Arg

Val Ser Arg Ile Phe Gln Cys Cys Ala Trp 70 75 Ser 65

22

Phe Ser Leu Leu Leu Phe Tyr Arg Val Phe Ile 다. 85 Gly Gly Val Phe

ž Leu Thr Ser Ile Phe 125 Pro Ser Asp Pro Val Leu Gln Ser Val Thr Ala Arg Ile Ile Gly 100 Trp Leu Glu Phe Phe 120 Gly Asp Val Trp Ser 115 His 8

Val Leu Pro Leu Phe Val Leu Ser Lys Val Val Asn 115 Ser Ala Leu Trp 35

Ala Ile Trp Phe Gln Asp Ile Ala Asp Leu Ala Phe Glu Val Ser Gly 145 Pro Phe Pro Ser Val Ser Lys lle lle Ala Asp Met 165 Arg Lys Pro His 6

Leu Phe Asn Leu Leu Gln Ala Leu Phe Leu Ile Gln Gly Met Phe 190 45

3 Ser Leu Leu Val 205 Ser Leu Phe Pro Ile His Leu Val Gly Gln 195 Val

Ser Leu Tyr Cys Phe Glu Tyr Arg Trp Phe 215 Met Ser Leu Leu Tyr 210 His S

Leu Thr Ala Asn Lys Gly Ile Glu Met His Gln Arg Leu Ser Asn Ile Glu Arg Asn 235 255 몺 Leu Ala Trp Pro Tyr Tyr Phe Gly Phe Gly Leu Pro 245 250

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Met Gln Ser Ser Tyr Ile Ile Ser Gly Cye Leu Phe Ser Ile Leu Phe $260\,$ 8

Pro Leu Phe Ile Ile Ser Ala Asn Glu Ala Lys Thr Pro Gly Lys Ala 275

Ser Asn Leu Val Val Phe Leu 300 Tyr Leu Phe Gln Leu Arg Leu Phe Ser 290

Arg Leu Phe His Lys Thr Val Tyr Leu Gln Ser Ala Leu Ser Ser 305 315

Thr Ser Ala Glu Lys Phe Pro Ser Pro His Pro Ser Pro Ala Lys Leu 325 335 2

Lys Ala Thr Ala Gly His 340 13

(2) INFORMATION FOR SEQ ID NO: 638;

8

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 529 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 638

22

Met Ala Lys Phe Met Thr Pro Val Ile Gln Asp Asn Pro Ser Gly Trp 1 5 10 15

Gly Pro Cys Ala Val Pro Glu Glu Gln Phe Arg Asp Met Pro Tyr Gln Pro 20 ഉ

Ser Ser Gln Phe Gly 60 Gly Ala Lys Val Ala Asp Trp Thr 45 Thr Asn Lys Tyr 95 5 ጟጜ Phe Ser Lys Gly Asp Arg Leu Thr Tyr Gln Asp Lys Arg 35

Phe 80 Ser Gly Gly Ser Gln Tyr Ala Tyr Phe His Glu Glu Asp Glu 65 6

Agn Gin Leu Val Asp Thư Ala Arg Thư Gin Lys Thư Ala Tyr Gin Arg 85 90

Asn Met Leu Gln Phe Asn Leu Gln Ile Leu Pro Lys Ser Ala Lys Gln 115 Arg Arg Arg Met Arg Phe Ala Gln Arg Asn Leu Arg Arg Asp Lys Asp $100 \ 100 \ 105$ 45

Gln Lys Lys Phe Gln Lys Gln Phe 140 <u>r</u>e Glu Arg Glu Arg Ile Arg 130 Ľys

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Phe Ser Gln Lys Pro Arg Asp Ser 155 Val Lys Glu Glu Met Asp 170 Trp Asp Gln Lys 150 ដូ ij Arg Ser Asp ' Val Arg Gln Lys Glu Val Val 55

Pro Gln Leu Met Lys Met Arg Tyr Leu Glu Val Ser Glu Pro Gln Asp 8

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6 8 35 30 25 20 15 45 5 Leu Gly Tyr Val Ser Arg Tyr His Val Lys Asp Ser Ser Arg His Val $450 \ \ 460$ Arg Gly Ala Val Ile Ala Thr Glu Leu Lys Asn Asn Ser Tyr Lys Leu 420 425 Asn Gly Glu Val 385 11e Glu Cys Cys Gly Ala Leu Glu Tyr Tyr Asp Lys Ala Phe Asp Arg 195 200 205 Phe Pro Asn Pro Asn Pro Phe Val Glu Asp Asp Met Asp Lys Asn Glu 340 345 Leu Thr Val Ser Glu Thr Ala Asn Glu Pro Pro Gln Asp Glu Gly Asn 290 295 300 Gly Ser Lys Leu Phe Phe Asp Lys Arg Asp Asn Ser Asp Phe Asp Leu 285 Ser Cys Thr Arg Ser Val Tyr Ser Trp Asp Ile Val Val Gln Arg Val 260 265 His Thr Val Thr Thr 225 Ile Thr Thr Arg Ser Glu Lys Pro Leu Arg Xaa Xaa Lys Arg Ile Phe 210 215 Ala Arg Trp Thr Cys Cys Ala Leu Leu Ala Gly Ser Glu Tyr Leu Lys 435 440 445 Ile Asp Leu Ile Val Arg Cys Glu His Asp Gly Val Met Thr Gly Ala 370 380 His Asn.Phe Ser Gln Gln Cys Leu Arg Met Gly Lys Glu Arg Tyr Asn 325 330 Thr Gln Gly Asn Val Phe Ala Thr Asp Ala Ile Leu Ala Thr Leu Met
245 250 Ser Arg His Cys Asn Gly Val Asp Trp Arg Gln Lys Leu Asp Ser Gln
410 415 Ile Ala Ser Val Ala Tyr Arg Tyr Arg Ser Gly Lys Leu Gly Asp Asp 355 Ser Phe Asn Ser Pro Arg Asn Leu Ala Met Glu Ala Thr Tyr Ile Asn 305 310 310 180 Ser Phe Ile Asn Ile Lys Thr Leu Asn Glu Trp Asp 390 395 Thr Asp Asp Pro Val Ile Arg Lys Leu Ala Lys 230 235 185

\$ S Š 8 ઝ 30 25 20 15 5 S His Phe Xaa Arg Val Gln Phe His Leu Lys Asn Phe Asp Met Val Ile 115 120 125 Val Tyx Lys Asp Tyx Sex Lys Lys Val Thr Met Ile Asn Ala Ile Pro 130 Ser Thr Ile Val Asp Asp Pro Glu Gly Phe Phe Glu Gln Gly Gly Trp Ser 180 Lys Tyr Thr Glu Gly Val Gln Ser Leu 165 Val Ala Ser Leu 145 Thr Glu Trp Pro Pro Phe Val Val Thr Leu Asp Glu Val Glu Leu Ile 100 110 Arg Ser Thr Cys Leu Gen Pro Thr Ser Ser Ala Leu Val Asn Ala 95 Glu Phe Glu Val Pro Phe Arg Asp Leu Gly Phe Asn Gly Ala Pro Tyr 65 70 75 Thr Thr Asp Leu Gly Lys His Gln His Met His Asp Arg Asp Asp Leu 20 25 Lys Lys Arg His Thr Asp Val Gln Phe Tyr Thr Glu Val Gly Glu Ile
1 5 10 15 Asn Lys Gln Val Ile Arg Val Tyr Ser Leu Pro Asp Gly Thr Phe 515 525 (2) INFORMATION FOR SEQ ID NO: 639: Ĭ, (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 639: 500 (A) LENGTH: 194 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Ąsp Pro Ile Lys Glu Trp Leu Asn Ser Cys Asp 150 155 505 Asn Trp Thr Lys Ile Met 170 510 160 Ser

(2) INFORMATION FOR SEQ ID NO: 640:

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Ile Leu Gly Thr Gln Gln Phe Lys Pro Asn Glu Phe Ala Ser Gln Ile 465 470 475

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Ile Cys Met Lys Leu Glu Glu Gly Lys Tyr Leu Ile Leu Lys Asp Pro

Asn Leu Ser Val Glu Asn Ala Trp Gly Ile Leu Arg Cys Val Ile Asp 485 , 490 495

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 640; (A) LENGTH: 70 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 3

Arg Ser Gly Leu Gly Leu Gly Ile Thr Ile Ala Phe Leu Ala Thr Leu

2

Ile Thr Gln Phe Leu Val Tyr Asn Gly Val Tyr Gln Tyr Thr Ser Pro 20 30

Asp Phe Leu Tyr 11e Arg Ser Trp Leu Pro Cys 11e Phe Ser Gly 45

2

Gly Val Thr Val Gly Asn Ile Gly Arg Gln Leu Ala Met Gly Val Pro 50 60

Glu Lys Pro His Ser Asp 65 70 2

(2) INFORMATION FOR SEQ ID NO: 641: 25

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 101 amino acids (B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 641: (D) TOPOLOGY: linear

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Val Thr Gin Pro Lys His Leu Ser Ala Ser Het Gly Gly Ser Val Glu 1 5 Ile Pro Phe Ser Phe Tyr Tyr Pro Trp Glu Leu Ala Xaa Xaa Pro Xaa 25 30

35

Val Arg Ile Ser Trp Arg Arg Gly His Phe His Gly Gln Ser Phe Tyr 45 **8**

Ser Thr Arg Pro Pro Ser Ile His Lys Asp Tyr Val Asn Arg Leu Phe 50 60

Leu Asn Try Thr Glu Gly Gln Glu Ser Gly Phe Leu Arg Ile Ser Asn 65 70 80 45

Leu Arg Lys Glu Asp Gin Ser Val Tyr Phe Cys Arg Val Glu Leu Asp 85

Thr Arg Arg Ser Gly 100

20

55

(2) INFORMATION FOR SEQ ID NO: 642:

LENGTH: 233 amino acids TYPE: amino acid (1) SEQUENCE CHARACTERISTICS: 3 9

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 642: (D) TOPOLOGY: linear

Met Glu Ala Gln Gln Val Asn Glu Ala Glu Ser Ala Arg Glu Gln Leu

Gln Xaa Leu His Asp Gln Ile Ala Gly Gln Lys Ala Ser Lys Gln Glu 20

Leu Glu Thr Glu Leu Glu Arg Leu Lys Gln Glu Phe His Tyr Ile Glu 18 \$452

Glu Asp Leu Tyr Arg Thr Lys Asn Thr Leu Gln Ser Arg Ile Lys Asp 50

15

Azg Asp Glu Glu Ile Gln Lys Leu Azg Asn Gln Leu Thr Asn Lys Thr 65 75 80

Leu Ser Asn Ser Gln Ser Glu Leu Glu Asn Arg Leu His Gln Leu 90 Thr Glu Thr Leu Ile Gln Lys Gln Thr Met Leu Glu Ser Leu Ser Thr 100

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Glu Lys Asn Ser Leu Val Phe Gln Leu Glu Arg Leu Glu Gln Gln Met 115

25

Asn Ser Ala Ser Gly Ser Ser Asn Gly Ser Ser Ile Asn Met Ser 110

Gly Ile Asp Asn Gly Glu Gly Thr Arg Leu Arg Asn Val Pro Val Leu 145 30

Phe Asn Asp Thr Glu Thr Asn Leu Ala Gly Met Tyr Gly Lys Val Arg

Lys Ala Ala Ser Ser Ile Asp Gln Phe Ser Ile Arg Leu Gly Ile Phe 180 33

Leu Arg Arg Tyr Pro Ile Ala Arg Val Phe Val Ile Ile Tyr Met Ala 195 Leu Leu His Leu Trp Val Met Ile Val Leu Leu Thr Tyr Thr Pro Glu 210

6

Met His His Asp Gln Pro Tyr Gly Lys 225

5

(2) INFORMATION FOR SEQ ID NO: 643

S

(A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (1) SEQUENCE CHARACTERISTICS:

55

Ile Arg His Glu Gln His Pro Asn Phe Ser Leu Glu Met His Ser Lys $_{\rm 1}$ 8

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 643

S

(2) INFORMATION FOR SEQ ID NO: 644:

5

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 63 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 644:

5

25 20 Thr Leu Thr Thr Cys Ser Gly Pro Thr Glu Lys Pro Ala Thr Lys Asn 35Arg His Pro Trp Val Ala Gly Ala Leu Val Gly Val Ser Gly Gly Leu 20

Tyr Phe Leu Lys Arg Leu Leu Gln Glu Met His Ile Arg Ala Asn $50\,$

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

Authorized officer Susan White PCT International Division	This sheet was received with the international application	E. SEPARATE FURNISHING O The indications listed below will be sub Number of Depart!")		D. DESIGNATED STATES FOR	C. ADDITIONAL INDICATIONS (team blank if not applicable)	Date of deposit February 26, 1997	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution Am	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to on page 116
Authorized officer	nemational application This sheet was received by the International Bureau on:	E. SEPARATE FURNISHING OF INDICATIONS (seaw blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Departs")	·	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)	NS (team blank if not applicable) This information is continued on an additional sheet	7 Accession Number 97897	uding postal code and country)	American Type Culture Collection	OSIT Further deposits are identified on an additional sheet	The indications made below relate to the microorganism referred to in the description on page 116 , line N/A

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 116	ed to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	lection
Address of depositary institution (including postal code and country)	(4
1230! Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209043
C. ADDITIONAL INDICATIONS (neave blank if not applicable)	(e) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIO	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (now blank if not applicable)	slank V not applicable)
The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	lureau later (specify the general nature of the indications, e.g., "Accession
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PCT/US98/04493

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ollection
Address of depositary institution (including postal code and country)	nuty)
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	."
Date of deposit September 4, 1997	Accession Number 209235
C. ADDITIONAL INDICATIONS (neave blank if not applicable)	able) This information is continued on an additional sheet
DESIGNATED STATES FOR WHICH INDICATIO	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE at the indications are and for all designated Secures.
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E. SEPAKATE FURNISHING OF INDICATIONS (name blank if not applicable)	s blank if not applicable)
he indications listed below will be submitted to the International umber of Deposit?	The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
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(PCT Rule 13bis)

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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IS ARE MADE (If the indications are not for all designated States)	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
e) This information is continued on an additional sheet	C. ADDITIONAL INDICATIONS (seem blank (f not applicable)
accession Number 97898	Date of deposit February 26, 1997
9)	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
ection	Name of depositary institution American Type Culture Collection
Further deposits are identified on an additional sheet 🔲	B. IDENTIFICATION OF DEPOSIT
d to in the description	A. The indications made below relate to the microorganism referred to in the description on page 122 , line N/A

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

Authorized officer Susan White PCT International Division	This sheet was received with the international application	For receiving Office use only	E. SEPARATE FURNISHING OF INDICATIONS (terre blank if not applicable) The indications listed below will be submitted to the international Bureau later (specify it humber of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARB MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit May 15, 1997	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page 122 , line N/A
Authorized officer	This sheet was received by the International Bureau on:	For International Bureau use only	E. SEPARATE FURNISHING OP INDICATIONS (tears blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	VS ARE MADE (If the indications are not for all designated States)	te) This information is continued on an additional sheet	Accession Number 209044	3)	lection	Further deposits are identified on an additional sheet	ed to in the description

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

	L	
A. The indications made below relate to the microorganism referred to in the description on page 126 , line N/A	-	A. The indications ma on page 126
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet	œi_	B. IDENTIFICATI
Name of depositary institution American Type Culture Collection	Pas V	Name of depositary ins
Address of depositary institution (including postal code and country)	Age	Address of depositary
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	L23 Roc Uni	12301 Parklawn Dri Rockville, Maryland United States of Am
Date of deposit February 26, 1997 Accession Number 97899	Date	Date of deposit May
C. ADDITIONAL INDICATIONS (name blant l'not applicable) This information is continued on an additional shect	Ü	ADDITIONAL I
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (() the indications are not for all designated States)	0.0	D. DESIGNATED S
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E. SEPARATE FURNISHING OF INDICATIONS (near blank if not applicable)	E. SI	E. SEPARATE FUR
The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	The in Numbe	The indications listed bel Number of Deposit")
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 126 (line N/A	red to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	llection
Address of depositary institution (including postal code and country)	(A)
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209045
C. ADDITIONAL INDICATIONS (teave blank if not applicable)	(e) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATION	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (neaw blank if not applicable)	slank (I not applicable)
The indications listed below will be submitted to the International E	The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

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lank () not applicable) wresu later (specify the general nature of the indications, e.g., "Accession	E. SEPARATE FURNISHING OF INDICATIONS (news blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
S ARE MADE (If the indications are not for all designated States)	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (1) the indications are not for all designated States)
e) This information is continued on an additional sheet 🔝	C. ADDITIONAL INDICATIONS (leave blank if not applicable)
Accession Number 209011	Date of deposit April 28, 1997
	12301 Parklawn Drive Rockville, Maryland 20852 United States of America
2	Address of depositary institution (including postal code and country)
ection	Name of depositary institution American Type Culture Collection
Further deposits are identified on an additional sheet 🔲	B. IDENTIFICATION OF DEPOSIT
d to in the description	A. The indications made below relate to the microorganism referred to in the description on page 130 , line N/A

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM (PCT Rule 13613)

For receiving Office use only ————————————————————————————————————	E. SEPARATE FURNISHING OF INDICATIONS (new blank if not applicable) The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (teams blank if not applicable) This information is continued on an additional sheet	Date of deposit February 26, 1997 Accession Number 97900	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT Funher deposits are identified on an additional sheet []	A. The indications made below relate to the microorganism referred to in the description on page 131 , line N/A
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Susan White PCT International Division

Authorized officer

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 137 , line N/A	re description	A. The indication
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	B. IDENTIFIC
Name of depositary institution American Type Culture Collection		Name of deposita
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America		Address of depos 12301 Parklawr Rockville, Mary United States of
Date of deposit February 26, 1997 Accession	Accession Number 97901	Date of deposit
C. ADDITIONAL INDICATIONS (new blank if not applicable) This	This information is continued on an additional sheet	C. ADDITION
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	MADE (If the indications are not for all designated States)	D. DESIGNAT
E. SEPARATE FURNISHING OF INDICATIONS Game hime if not amplicables	milicakisi	E. SEPARATE
The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")	t (specify the general nature of the indications, e.g., "Accession	The indications list Number of Deposit")
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 131 , time N/A	red to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ilection
Address of depositary institution (including postal code and country)	(61)
12301 Parkawn Drive Rockville, Maryland 20852 United States of America	
Datc of deposit May 15, 1997	Accession Number 209046
C. ADDITIONAL INDICATIONS (teave blank y not applicable)	ble) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIO	 DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (new blank if not applicable)	blank (f not applicable)
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 136is)

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NS ARE MADE (If the indications are not for all designated States)	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
le) This information is continued on an additional sheet 🔲	C. ADDITIONAL INDICATIONS (leave blank if not applicable)
Accession Number 209047	Date of deposit May 15, 1997
7)	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
lection	Name of depositary institution American Type Culture Collection
Further deposits are identified on an additional sheet 📋	B. IDENTIFICATION OF DEPOSIT
ed to in the description	A. The indications made below relate to the microorganism referred to in the description on page 137 , line N/A

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM
(PCT Rule 13bis)

Name of depositary institution American Type Culture Collection Address of depositary institution (including postal code and country) 1230 Parklawn Drive Rockville, Manyland 20852 United States of America C. ADDITIONAL INDICATIONS (new block (frost applicable)) Date of deposit May 22, 1997 C. ADDITIONAL INDICATIONS (new block (frost applicable)) D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (frost applicable) E. SEPARATE FURNISHING OF INDICATIONS (new block (frost applicable)) For receiving Office use only	Authorized officer Sugan White
American Type Culture Collection (including postal code and country) (including postal code and country) Accession Number 209076 [1997] Accession Number 209076 [CATIONS (Reave blank (I not applicable)] This information is continued on an additional sheet [CATIONS (Reave blank (I not applicable)]] FES FOR WHICH INDICATIONS (Reave blank (I not applicable)] Will be submitted to the International Bureau later (speelf) the general nature of the Indications, e.g., "Accessing Office use only	This sheet was received with the international application
n American Type Culture Collection (including postal code and country) 1997 Accession Number 209076 CATIONS (term blank l/not applicable) This information is continued on an additional sheet CATIONS (term blank l/not applicable) This information is continued on an additional sheet CATIONS (term blank l/not applicable) This information is continued on an additional sheet State. TES FOR WHICH INDICATIONS (term blank l/not applicable) Will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number 209076	For receiving Office use only
TES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated State.	EPARATE FURNISHING OF INDICATIONS indications listed below will be submitted to the Interna of Depati ^e 7
n American Type Culture Collection 1100 (including postal code and country) 12 1997 Accession Number 209076 CATIONS (leaw blank (I not applicable) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable)) This information is continued on an additional sheet CATIONS (Pleaw blank (I not applicable))	
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American Type Culture Collection American Type Culture Collection (including postal code and country) 2 Accession Number 209076 Accession Number 209076	·
FDEPOSIT Further deposits are identified on an additional sheet American Type Culture Collection (including postal code and country) 2 Accession Number 209076	C. ADDITIONAL INDICATIONS (team blank if not applicable)
FDEPOSIT Further deposits are identified on an additional sheet American Type Culture Collection Including postal code and country)	Date of deposit May 22, 1997
F DEPOSIT Further deposits are identified on an additional sheet American Type Culture Collection (including postal code and country)	12301 Parklawn Drive Rockville, Maryland 20852 United States of America
F DEPOSIT Further deposits are identified on an additional sheet American Type Culture Collection	Address of depositary institution (including postal code and country)
Further deposits are identified on an additional sheet	Name of depositary institution American Type Cultur
	IDENTIFICATION OF DEPOSIT
A. The indications made below relate to the microorganism referred to in the description on page 137 . line N/A	The indications made below relate to the microorganism on page 137 , line

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

	(ref Rule 13013)	
A. The indications made below relate to the microorganism referred to in the description on page 140	ned to in the description A	A. The indication: on page 16
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet.	B. IDENTIFICA
Name of depositary institution American Type Culture Collection	illection	Name of depositary
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	(12)	Address of deposit 12301 Parklawn Rockville, Maryl United States of
Date of deposit August 21, 1997	Accession Number 209215	Date of deposit
C. ADDITIONAL INDICATIONS from blank if not applicable)	bby This information is continued on an additional sheet	C. ADDITIONA
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	NS ARE MADE (if the indications are not for all designated States)	D. DESIGNATE
E. SEPARATE FURNISHING OF INDICATIONS (terre blank l/not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Depaul")	blank l'Inot applicable) Jureau later (specify the general nature of the Indications, e.g., "Accession	E. SEPARATE F The indications lister Number of Deposit 7
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PCT/US98/04493

A. The indications made below relate to the microorganism referred to in the description on page 160 , line N/A	rred to in the description A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary Institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	игу)
Date of deposil February 26, 1997	Accession Number 97904
C. ADDITIONAL INDICATIONS (new blank if not applicable)	able) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATION	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (name blank if not applicable)	: blank (f not applicable)
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., Number of Deposit")	Bureau later (specify the general nature of the indications, e.g., "Accession
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 154 , line N/A	d to in the description
B. IDENTIFICATION OF DEPOSIT	Funher deposits are identified on an additional sheet
Name of depositary institution . American Type Culture Collection	ection
Address of depositary institution (including postal code and country)	
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit July 3, 1997	Accession Number 209139
C. ADDITIONAL INDICATIONS (teave blank if not applicable)	e) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	S ARE MADE (If the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (trave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify it	viank () not applicable) ureau later (specify the general nature of the indications, e.g., "Accession
The indications listed below will be submitted to the International Bureau later (pecify the general nature of the indications, e.g., "Accession Number of Deposit")	ureau later (specify the general nature of the indications, e.g., "Accession
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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(PCT Rule 136is)

For receiving Office use onlyFor International Bureau use only	E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accessions Humber of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (() the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (theore blank if not applicable) This information is continued on an additional sheet	Date of deposit May 15, 1997 Accession Number 209049	Address of depositary institution (including postal code and country) [230] Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet	A. The indications made below relate to the microorganism referred to in the description on page 153 , line N/A

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Susan White PCT International Division

Authorized officer

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page $$153$$, line $$\mathrm{N/A}$$	cd to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	lection
Address of depositary institution (including postal code and country)	. (4
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit February 26, 1997	Accession Number 97903
C. ADDITIONAL INDICATIONS (team blank if not applicable)	(4) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	IS ARE MADE (If the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (new blank if not applicable)	ılank if not applicable)
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Authorized officer Susan White PCT international Division	Authorized officer

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

A. The indications made below relate to the microorganism referred to in the description on page 142 . Inc N/A	ed to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	lection
Address of depository institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	(£
Date of deposit June 12, 1997	Accession Number 209119
C. ADDITIONAL INDICATIONS (teave blank if not applicable)	1e) This information is continued on an additional sheet
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

Authorized officer Susain Willie Authorized officer PCT Internetional Division	For receiving Office use only For International Bureau use only For International Bureau use only This sheet was received by the International Bureau on:	E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (teave blank if not applicable) This information is continued on an additional sheet	coession Number 97902	Address of depositary institution (<i>including postal code and country</i>) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America		3	A. The indications made below relate to the microorganism referred to in the description on page 146 , line N/A
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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Authorized officer Sussan White PCT International Division	This sheet was received with the international application	Ear manisting Office use only	E. SEPARATE FURNISHING OF INDICATIONS sterve blank if not applicable) The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession number of Depart")		D. DESIGNATED STATES FOR WHICH INDICATION	C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit May 15, 1997	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page 146 , line N/A
Authorized officer	This sheet was received by the International Bureau on:	For International Bureau use only	iank (I not applicable) wreas later (specify the general nature of the indications, e.g., "Accession	·	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)	This information is continued on an additional sneet	Accession Number 209048		tion	Further deposits are identified on an additional sheet	to in the description

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A. The indications made below relate to the microorganism referred to in the description	on page 160 . line N/A B. IDENTIFICATION OF DEPOSIT	Name of depositary institution American Type Culture Collection	Address of depositury institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Date of deposit May 15, 1997	C. ADDITIONAL INDICATIONS (team blank l) not applicable)	D. DESIGNATED STATES FOR WHICH INDICATIO	E. SEPARATE FURNISHING OF INDICATIONS (teaw blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	For receiving Office use only	This sheet was received with the international application	Authorized officer Susan White PCT International Division

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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

A. The indications made below relate to the microorganism referred to in the description on page 142 , line N/A	eferred to in the description N/A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet 🔯
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including pastal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	(Сии
Date of deposit February 12, 1998	Accession Number 209627
C. ADDITIONAL INDICATIONS (seaw blank if not applicable)	able) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIC	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (grine indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (naw blank i) not applicable)	blank if not applicable)
The indications listed below will be submitted to the International Vumber of Deposit 7	The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
For receiving Office use only	For International Bureau use only
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Form PCT/RC/114 (Islv 1902)	

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What Is Claimed Is:

nucleotide sequence at least 95% identical to a sequence selected from the group An isolated nucleic acid molecule comprising a polynucleotide having a

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- the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of
- polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X; (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a

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- which is hybridizable to SEQ ID NO:X; polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a
- which is hybridizable to SEQ ID NO:X; polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a

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- sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA
- having biological activity;

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- (f) a polynucleotide which is a variant of SEQ ID NO:X;
- (g) a polynucleotide which is an allelic variant of SEQ ID NO:X
- (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
- (i) a polynucleotide capable of hybridizing under stringent conditions to any
- 25 one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not sequence of only A residues or of only T residues. hybridize under stringent conditions to a nucleic acid molecule having a nucleotide
- 30 polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein. The isolated nucleic acid molecule of claim 1, wherein the
- in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X. polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included The isolated nucleic acid molecule of claim 1, wherein the

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- polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID The isolated nucleic acid molecule of claim 1, wherein the
- sequence comprises sequential nucleotide deletions from either the C-terminus or the Nterminus. The isolated nucleic acid molecule of claim 2, wherein the nucleotide
- 5 sequence comprises sequential nucleotide deletions from either the C-terminus or the Nterminus. The isolated nucleic acid molecule of claim 3, wherein the nucleotide
- claim 1. A recombinant vector comprising the isolated nucleic acid molecule of

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- nucleic acid molecule of claim 1. A method of making a recombinant host cell comprising the isolated
- ঞ A recombinant host cell produced by the method of claim 8

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- 10. The recombinant host cell of claim 9 comprising vector sequences.
- identical to a sequence selected from the group consisting of: An isolated polypeptide comprising an amino acid sequence at least 95%

25

- (a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included
- in ATCC Deposit No:Z;
- in ATCC Deposit No:Z, having biological activity; (b) a polypeptide fragment of SEQ ID NO: Y or the encoded sequence included
- ၓ ATCC Deposit No:Z; (c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in
- ATCC Deposit No:2; (d) a polypeptide epitope of SEQ ID NO: Y or the encoded sequence included in
- ATCC Deposit No:Z; (e) a secreted form of SEQ ID NO:Y or the encoded sequence included in

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ATCC Deposit No:Z; (f) a full length protein of SEQ ID NO:Y or the encoded sequence included in

- (g) a variant of SEQ ID NO:Y;
- (h) an allelic variant of SEQ ID NO:Y; or
- (i) a species homologue of the SEQ ID NO:Y.
- The isolated polypeptide of claim 11, wherein the secreted form or the
 - 'n comprises sequential amino acid deletions from either the C-terminus full len or th

'body that binds specifically to the isolated polypeptide of

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*xpresses the isolated polypeptide of claim

7.

A method.

· comprising:

under conditions such that

said polypeptide is expressed; and (a) culturing the recon.

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(b) recovering said polypeptide.

The polypeptide produced by claim 15. 16.

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comprising administering to a mammalian subject a therapeutically effective amount of A method for preventing, treating, or ameliorating a medical condition, the polypeptide of claim 11 or the polynucleotide of claim 1. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising: . 8 23

(a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and

(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation. 39

- A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:
- (a) determining the presence or amount of expression of the polypeptide of
 - claim 11 in a biological sample; and 35
- (b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide

A method for identifying a binding partner to the polypeptide of claim 11

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comprising:

(a) contacting the polypeptide of claim 11 with a binding partner; and

(b) determining whether the binding partner effects an activity of the polypeptide. The gene corresponding to the cDNA sequence of SEQ ID NO Y.

A method of identifying an activity in a biological assay, wherein the method comprises: 22.

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(a) expressing SEQ ID NO:X in a cell;

(b) isolating the supernatant;

(c) detecting an activity in a biological assay; and

(d) identifying the protein in the supernatant having the activity.

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The product produced by the method of claim 22. 23.